## GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 5, 2005, 15:38:47; Search time 8644.89 Seconds

(without alignments)

12117.322 Million cell updates/sec

Title: US-10-624-932-1

Perfect score: 2752

Sequence: 1 ccgcggggcccggcccgg.....tgagtgctgaggccggccag 2752

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 34239544 segs, 19032134700 residues

Total number of hits satisfying chosen parameters: 68479088

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : EST:\*

1: gb\_est1:\*

2: gb\_est2:\*

3: gb htc:\*

4: gb est3:\*

5: gb est4:\*

6: gb est5:\*

7: gb\_est6:\*

8: gb gss1:\*

9: gb\_gss2:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

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R	esult		Query				
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	2	950.4	34.5	2791	9	AY406493	AY406493 Mus muscu
	3	923.4	33.6	3790	3	AK031655	AK031655 Mus muscu
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	5	872.4	31.7	3866	3	AK018177	AK018177 Mus muscu
	6	814	29.6	2802	9	AY406492	AY406492 Pan trogl
	7	810.4	29.4	2532	9	AY411747	AY411747 Homo sapi
	8	780.4	28.4	2532	9	AY411749	AY411749 Mus muscu

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VERSION
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KEYWORDS
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            Homo sapiens (human)
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REFERENCE
            1 (bases 1 to 2802)
  AUTHORS
            Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A.,
            Todd, M.A., Tanenbaum, D.M., Civello, D.R., Lu, F., Murphy, B.,
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RESULT 1

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Ferriera, S., Wang, G., Zheng, X.H., White, T.J., Sninsky, J.J.,
          Adams, M.D. and Cargill, M.
          Inferring nonneutral evolution from human-chimp-mouse orthologous
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          Adams, M.D. and Cargill, M.
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          Submitted (16-NOV-2003) Celera Genomics, 45 West Gude Drive,
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          This sequence was made by sequencing genomic exons and ordering
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                                                         . | | | | | | |
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VERSION AY406493.1 GI:39762467 KEYWORDS GSS. SOURCE Mus musculus (house mouse) ORGANISM Mus musculus

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1 (bases 1 to 2791) REFERENCE

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Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A.,
 AUTHORS
          Todd, M.A., Tanenbaum, D.M., Civello, D.R., Lu, F., Murphy, B.,
           Ferriera, S., Wang, G., Zheng, X.H., White, T.J., Sninsky, J.J.,
          Adams, M.D. and Cargill, M.
 TITLE
           Inferring nonneutral evolution from human-chimp-mouse orthologous
           gene trios
          Science 302 (5652), 1960-1963 (2003)
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  PUBMED
           14671302
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REFERENCE
 AUTHORS
          Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A.,
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          Adams, M.D. and Cargill, M.
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 TITLE
 JOURNAL
           Submitted (16-NOV-2003) Celera Genomics, 45 West Gude Drive,
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SOURCE ORGANI		Mus musculus (house mouse) Mus musculus
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REFERENC AUTHOR	E	Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus. 1 Carninci, P. and Hayashizaki, Y.
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High-efficiency full-length cDNA cloning
  TITLE
            Meth. Enzymol. 303, 19-44 (1999)
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REFERENCE
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            Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
 AUTHORS
            Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
            Normalization and subtraction of cap-trapper-selected cDNAs to
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            prepare full-length cDNA libraries for rapid discovery of new genes
            Genome Res. 10 (10), 1617-1630 (2000)
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            Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P.,
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            RIKEN integrated sequence analysis (RISA) system--384-format
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            sequencing pipeline with 384 multicapillary sequencer
            Genome Res. 10 (11), 1757-1771 (2000)
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            20530913
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            11076861
REFERENCE
            The RIKEN Genome Exploration Research Group Phase II Team and the
  AUTHORS
            FANTOM Consortium.
            Functional annotation of a full-length mouse cDNA collection
  TITLE
  JOURNAL
            Nature 409, 685-690 (2001)
REFERENCE
            5
            The FANTOM Consortium and the RIKEN Genome Exploration Research
  AUTHORS
            Group Phase I & II Team.
  TITLE
            Analysis of the mouse transcriptome based on functional annotation
            of 60,770 full-length cDNAs
            Nature 420, 563-573 (2002)
  JOURNAL
            6 (bases 1 to 3790)
REFERENCE
            Adachi, J., Aizawa, K., Akimura, T., Arakawa, T., Bono, H., Carninci, P.,
  AUTHORS
            Fukuda, S., Furuno, M., Hanagaki, T., Hara, A., Hashizume, W.,
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  TITLE
            Direct Submission
            Submitted (16-JUL-2001) Yoshihide Hayashizaki, The Institute of
  JOURNAL
            Physical and Chemical Research (RIKEN), Laboratory for Genome
            Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
            RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
            Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.jp,
            URL: http://genome.gsc.riken.jp/, Tel:81-45-503-9222,
            Fax: 81-45-503-9216)
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COMMENT
           cDNA library was prepared and sequenced in Mouse Genome
           Encyclopedia Project of Genome Exploration Research Group in Riken
           Genomic Sciences Center and Genome Science Laboratory in RIKEN.
           Division of Experimental Animal Research in Riken contributed to
           prepare mouse tissues.
           Please visit our web site for further details.
           URL:http://genome.gsc.riken.jp/
           URL:http://fantom.gsc.riken.jp/.
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 AUTHORS
           Li, W.B., Gruber, C., Jessee, J. and Polayes, D.
           Full-length cDNA libraries and normalization
  TITLE
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           Unpublished
  REMARK
           Contact : Feng Liang Email : fliang@lifetech.com URL :
           http://fulllength.invitrogen.com/ InVitroGen Corporation 1600
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REFERENCE
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 AUTHORS
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           - Web : www.genoscope.cns.fr)
COMMENT
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REFERENCE
  AUTHORS
            Carninci, P. and Hayashizaki, Y.
            High-efficiency full-length cDNA cloning
  TITLE
            Meth. Enzymol. 303, 19-44 (1999)
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            Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K.,
  AUTHORS
            Itoh, M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
            Normalization and subtraction of cap-trapper-selected cDNAs to
  TITLE
            prepare full-length cDNA libraries for rapid discovery of new genes
            Genome Res. 10 (10), 1617-1630 (2000)
  JOURNAL
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REFERENCE
            Shibata, K., Itoh, M., Aizawa, K., Nagaoka, S., Sasaki, N., Carninci, P.,
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            Konno, H., Akiyama, J., Nishi, K., Kitsunai, T., Tashiro, H., Itoh, M.,
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  TITLE
            RIKEN integrated sequence analysis (RISA) system--384-format
            sequencing pipeline with 384 multicapillary sequencer
            Genome Res. 10 (11), 1757-1771 (2000)
  JOURNAL
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REFERENCE
            The RİKEN Genome Exploration Research Group Phase II Team and the
  AUTHORS
            FANTOM Consortium.
            Functional annotation of a full-length mouse cDNA collection
  TITLE
  JOURNAL
            Nature 409, 685-690 (2001)
REFERENCE
  AUTHORS
            The FANTOM Consortium and the RIKEN Genome Exploration Research
            Group Phase I & II Team.
  TITLE
            Analysis of the mouse transcriptome based on functional annotation
            of 60,770 full-length cDNAs
  JOURNAL
            Nature 420, 563-573 (2002)
REFERENCE
                (bases 1 to 3866)
            Adachi, J., Aizawa, K., Akahira, S., Akimura, T., Arai, A., Aono, H.,
  AUTHORS
            Arakawa, T., Bono, H., Carninci, P., Fukuda, S., Fukunishi, Y.,
            Furuno, M., Hanagaki, T., Hara, A., Hayatsu, N., Hiramoto, K.,
            Hiraoka, T., Hori, F., Imotani, K., Ishii, Y., Itoh, M., Izawa, M.,
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Saito, H., Saito, R., Sakai, C., Sakai, K., Sano, H., Sasaki, D.,
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      Tejima, Y., Toya, T., Yamamura, T., Yasunishi, A., Yoshida, K.,
      Yoshino, M., Muramatsu, M. and Hayashizaki, Y.
      Direct Submission
      Submitted (10-JUL-2000) Yoshihide Hayashizaki, The Institute of
      Physical and Chemical Research (RIKEN), Laboratory for Genome
      Exploration Research Group, RIKEN Genomic Sciences Center (GSC),
      RIKEN Yokohama Institute; 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama,
      Kanagawa 230-0045, Japan (E-mail:genome-res@gsc.riken.jp,
      URL: http://genome.gsc.riken.jp/, Tel:81-45-503-9222,
      Fax:81-45-503-9216)
      Please visit our web site (http://genome.gsc.riken.jp/) for further
      details.
      cDNA library was prepared and sequenced in Mouse Genome
      Encyclopedia Project of Genome Exploration Research Group in Riken
      Genomic Sciences Center and Genome Science Laboratory in RIKEN.
      Division of Experimental Animal Research in Riken contributed to
      prepare mouse tissues. First strand cDNA was primed with a primer
      prepared by using trehalose thermo-activated reverse transcriptase
      and subsequently enriched for full-length by cap-trapper. cDNA went
      through one round of normalization to Rot = 10.0 and subtraction to
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      with BamHI and XhoI. Vector: a modified pBluescript KS(+) after
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TITLE

COMMENT

**FEATURES** 

CDS

JOURNAL

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## ORIGIN

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           Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A.,
  AUTHORS
           Todd, M.A., Tanenbaum, D.M., Civello, D.R., Lu, F., Murphy, B.,
           Ferriera, S., Wang, G., Zheng, X.H., White, T.J., Sninsky, J.J.,
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  TITLE
           Inferring nonneutral evolution from human-chimp-mouse orthologous
           Science 302 (5652), 1960-1963 (2003)
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           Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A.,
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Submitted (16-NOV-2003) Celera Genomics, 45 West Gude Drive,

**JOURNAL** 

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          Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A.,
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Direct Submission
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   JOURNAL
                   Submitted (16-NOV-2003) Celera Genomics, 45 West Gude Drive,
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COMMENT
                   This sequence was made by sequencing genomic exons and ordering
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Adams, M.D. and Cargill, M.

Qv

11 111

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           Direct Submission
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 JOURNAL
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 AUTHORS
           NIH-MGC http://mgc.nci.nih.gov/.
           National Institutes of Health, Mammalian Gene Collection (MGC)
 TITLE
 JOURNAL
           Unpublished (1999)
           Contact: Robert Strausberg, Ph.D.
COMMENT
           Email: cgapbs-r@mail.nih.gov
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            cDNA Library Preparation: Life Technologies, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Incyte Genomics, Inc.
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           found through the I.M.A.G.E. Consortium/LLNL at:
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  AUTHORS
            Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G.,
            Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, C.M., Schuler, G.D.,
            Altschul, S.F., Zeeberg, B., Buetow, K.H., Schaefer, C.F., Bhat, N.K.,
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            Diatchenko, L., Marusina, K., Farmer, A.A., Rubin, G.M., Hong, L.,
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            Butterfield, Y.S., Krzywinski, M.I., Skalska, U., Smailus, D.E.,
            Schnerch, A., Schein, J.E., Jones, S.J. and Marra, M.A.
  TITLE
            Generation and initial analysis of more than 15,000 full-length
            human and mouse cDNA sequences
            Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
  JOURNAL
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  AUTHORS
            Strausberg, R.
  TITLE
            Direct Submission
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            Submitted (02-JUL-2002) National Institutes of Health, Mammalian
            Gene Collection (MGC), Cancer Genomics Office, National Cancer
            Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
            NIH-MGC Project URL: http://mgc.nci.nih.gov
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COMMENT
            Contact: MGC help desk
            Email: cgapbs-r@mail.nih.gov
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            cDNA Library Preparation: Life Technologies, Inc.
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            DNA Sequencing by: National Institutes of Health Intramural
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Sequencing Center (NISC),

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Contact: nisc mgc@nhgri.nih.gov
          Akhter, N., Ayele, K., Beckstrom-Sternberg, S.M., Benjamin, B.,
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Gaithersburg, Maryland;

Web site: http://www.nisc.nih.gov/

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              (bases 1 to 788)
REFERENCE
           NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 AUTHORS
 TITLE
           National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
           Tumor Gene Index
           Unpublished (1997)
  JOURNAL
           Contact: Robert Strausberg, Ph.D.
COMMENT
           Email: cgapbs-r@mail.nih.gov
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           R. Emmert-Buck, M.D., Ph.D. cDNA Library Preparation: M. Bento
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                    strand cDNA was primed with a Not I - oligo(dT) primer [5'
                    double-stranded cDNA was ligated to Eco RI adaptors
                    (Pharmacia), digested with Not I and cloned into the Not
                    I and Eco RI sites of the modified pT7T3 vector. Library
                    went through one round of normalization, and was
                    constructed by Bento Soares and M. Fatima Bonaldo.
ORIGIN
                        26.7%;
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Qv

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Db	421	ACTACTGCCAGCTGGAGGCCAGTGCCTACGTCTTCACCGAGCAGCTGAGCCGCTATG	480
Qу	1949	CCCTGGTGGGAGAGGCCCTCAGCGTGGCTGCCGCCAAGCGCCTCAAGCTGCTTCTGTTTG	2008
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LOCUS BX348193 796 bp mRNA linear EST 08-APR-2004 DEFINITION BX348193 Homo sapiens NEUROBLASTOMA COT 10-NORMALIZED Homo sapiens

cDNA clone CS0DB008YE02 5-PRIME, mRNA sequence.

ACCESSION BX348193

VERSION BX348193.2 GI:46286231

KEYWORDS EST.

SOURCE Homo sapiens (human)

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ORGANISM
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          Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
          Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
             (bases 1 to 796)
REFERENCE
          Li, W.B., Gruber, C., Jessee, J. and Polayes, D.
 AUTHORS
          Full-length cDNA libraries and normalization
 TITLE
          Unpublished (2001)
 JOURNAL
          On May 5, 2003 this sequence version replaced gi:30367258.
COMMENT
          Contact: Genoscope
          Genoscope - Centre National de Sequencage
          2 rue Gaston Cremieux, CP 5706 - 91057 EVRY cedex - FRANCE
          Email: seqref@genoscope.cns.fr, Web : www.genoscope.cns.fr
          1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime
          end enriched, double-strand cDNA was digested with Not I and cloned
          into the Not I and EcoR V sites of the pCMVSPORT 6 vector. Library
          was normalized. Library was constructed by Life Technologies, a
          division of Invitrogen. This sequence belongs to sequence cluster
          3239.r
          For more information about this cluster, see
          http://www.genoscope.cns.fr/cdna?s=CSOBAF004ZD01 AF00293 1&c=3239.r
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TITLE JOURNAL COMMENT	Na Ui	National Institutes of Health, Mammalian Gene Collection (MGC) Unpublished (1999)								
COLTABINI		ontact: Robert Strausberg, Ph.D. mail: cgapbs-r@mail.nih.gov								

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Tissue Procurement: Life Technologies, Inc.
           cDNA Library Preparation: Life Technologies, Inc.
           cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
           DNA Sequencing by: Incyte Genomics, Inc.
           Clone distribution: MGC clone distribution information can be
          found through the I.M.A.G.E. Consortium/LLNL at:
          http://image.llnl.gov
          Plate: LLAM11434 row: 1 column: 16
          High quality sequence stop: 744.
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                  source anonymous pool of 6 male brains, age range 23-27; 1
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                  oligo-dT primed and directionally cloned (EcoRV site is
                  destroyed upon cloning). Average insert size 1.8 kb,
                  insert size range 1-3 kb. Library is normalized and
                  enriched for full-length clones and was constructed by C.
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                  021. Note: this is a NIH MGC Library."
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Qу

Db

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           AY411748
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  AUTHORS
           Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A.,
           Todd, M.A., Tanenbaum, D.M., Civello, D.R., Lu, F., Murphy, B.,
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 TITLE
           Inferring nonneutral evolution from human-chimp-mouse orthologous
           gene trios
           Science 302 (5652), 1960-1963 (2003)
  JOURNAL
  PUBMED
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REFERENCE
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 AUTHORS
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           Ferriera, S., Wang, G., Zheng, X.H., White, T.J., Sninsky, J.J.,
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           Direct Submission
  TITLE
  JOURNAL
           Submitted (16-NOV-2003) Celera Genomics, 45 West Gude Drive,
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           This sequence was made by sequencing genomic exons and ordering
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Job time: 8663.89 secs

## GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 5, 2005, 15:11:26; Search time 11689.8 Seconds

(without alignments)

11407.261 Million cell updates/sec

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Gapop 10.0 , Gapext 1.0

Searched: 4708233 segs, 24227607955 residues

Total number of hits satisfying chosen parameters: 9416466

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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13: gb\_un:\*

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

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 AUTHORS
        Padigaru, M., Mezes, P., Mishra, V., Burgess, C., Casman, S.,
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Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

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Db		GTGAACGGT	GGGTGGTC	SACGT	'GGAC	CGAG1	GGTC	CGTCT	GCAGCG	CCAGCTG	TGGGCG	C 780
Qу	020											

Db	781	GGCTGGCAGAAACGGAGCCGGAGCTGCACCAACCCGGCGCCTCTCAACGGGGGCGCTTTC	840
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Qу	946	TGGAGCCCGTGGAGCAAGTGGTCGGCCTGTGGGCTGGACTGCACCCACTGGCGGAGCCGT	1005
Db	901	TGGAGCCCGTGGAGCAAGTGGTCGGCCTGTGGGCTGGACTGCACCCACTGGCGGAGCCGT	960
Qу	1006	GAGTGCTCTGACCCAGCACCCCGCAACGGAGGGAGGAGGAGTGCCAGGGCACTGACCTGGAC	1065
Db	961	GAGTGCTCTGACCCAGCACCCCGCAACGGAGGGAGGGAGG	1020
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Db	1021	ACCCGCAACTGTACCAGTGACCTCTGTGTACACACTGCTTCTGGCCCTGAGGACGTGGCC	1080
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Qy	1246	ACCTCAGGCTTCCAGCCCGTCAGCATCAAGCCCAGCAAAGCAGACAACCCCCATCTGCTC	1305
Db	1201	ACCTCAGGCTTCCAGCCCGTCAGCATCAAGCCCAGCAAAGCAGACAACCCCCATCTGCTC	1260
Qу	1306	ACCATCCAGCCGGACCTCAGCACCACCACCACCACCAGGGCAGTCTCTGTCCCCGG	1365
Db	1261	ACCATCCAGCCGGACCTCAGCACCACCACCACCACCAGGGCAGTCTCTGTCCCCGG	1320
QУ	1366	CAGGATGGGCCCAGCCCCAAGTTCCAGCTCACCAATGGGCACCTGCTCAGCCCCCTGGGT	1425
Db	1321	CAGGATGGGCCCAGCCCCAAGTTCCAGCTCACCAATGGGCACCTGCTCAGCCCCCTGGGT	1380
Qу	1426	GGCGGCCGCCACACTGCACCACAGCTCTCCCACCTCTGAGGCCGAGGAGTTCGTCTCC	1485
Db	1381	GGCGGCCGCCACACTGCACCACACTCTCCCACCTCTGAGGCCGAGGAGTTCGTCTCC	1440
Qу	1486	CGCCTCTCCACCCAGAACTACTTCCGCTCCCTGCCCCGAGGCACCAGCAACATGACCTAT	1545
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QУ	1546	GGGACCTTCAACTTCCTCGGGGGCCGGCTGATGATCCCTAATACAGGTATCAGCCTCCTC	1605
Db	1501	GGGACCTTCAACTTCCTCGGGGGCCGGCTGATGATCCCTAATACAGGAATCAGCCTCCTC	1560
Qу	1606	ATCCCCCAGATGCCATACCCCGAGGGAAGATCTATGAGATCTACCTCACGCTGCACAAG	1665
Db	1561	ATCCCCCAGATGCCATACCCCGAGGGAAGATCTATGAGATCTACCTCACGCTGCACAAG	1620
Qу	1666	CCGGAAGACGTGAGGTTGCCCCTAGCTGGCTGTCAGACCCTGCTGAGTCCCATCGTTAGC	1725
Db	1621	CCGGAAGACGTGAGGTTGCCCCTAGCTGGCTGTCAGACCCTGCTGAGTCCCATCGTTAGC	1680

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Db	1681	TGTGGACCCCTGGCGTCCTGCTCACCCGGCCAGTCATCCTGGCTATGGACCACTGTGGG	1740
Qy	1786	GAGCCCAGCCCTGACAGCTGGAGCCTGCGCCTCAAAAAGCAGTCGTGCGAGGGCAGCTGG	1845
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Qy	1846	GAGGATGTGCTGCACCTGGGCGAGGAGGCGCCCTCCCACCTCTACTACTGCCAGCTGGAG	1905
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Qy	2266	CAGCGGTACTTGCACTGCACCTTCACCCTGGAGCGTGTCAGCCCCAGCACTAGTGACCTG	2325
Db	2221	CAGCGGTACTTGCACTGCACCTTCACCCTGGAGCGTGTCAGCCCCAGCACTAGTGACCTG	2280
Qy	2326	GCCTGCAAGCTGTGGGTGTGGCAGGTGGAGGGCGACGGGCAGAGCTTCAGCATCAACTTC	2385
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Db	2341		2400
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Db	2401		2460
Qу	2506	TCCAGCCTGGACCCACCCTGTAGGCGGGGTGCCGACTGGCGGACTCTGGCCCAGAAACTC	2565
Db	2461	TCCAGCCTGGACCCACCCTGTAGGCGGGGTGCCGACTGGCGGACTCTGGCCCAGAACTC	2520

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Qy	2626 AACCTGTGGGAGGCGCGCACTTCCCCAACGGCAACCTCAGCCAGC
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Qy	2686 GCTGGACTGGCCAGCCAGACGCTGGCCTCTTCACAGTGTCGGAGGCTGAGTGCTGA 2742
Db	2641 GCTGGACTGGCCAGACGCTGGCCTCTTCACAGTGTCGGAGGCTGAGTGCTGA 2697
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Db	
Qу	181 TTCCTGGTGGAGCCCGAGGATGTGTACATCGTCAAGAACAAGCCAGTGCTGCTTGTGTGC 240
Db	

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DЪ	342	GACCACGTGATCGAGCGCAGCACAGACGGGAGCAGTGGTGAGCCGACCATGGAGGTCCGC	401
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Qу	541	ATCGTGCTGCCCTCCACCGGAGGGCATCCCTCCAGCCGAGGTGGAGTGGCTCCGG	600
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Qу	1258	CAGCCCGTCAGCATCAAGCCCAGCAAAGCAGACAACCCCCATCTGCTCACCATCCAGCCG	1317
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Qу	1498	CAGAACTACTTCCGCTCCCTGCCCCGAGGCACCAGCAACATGACCTATGGGACCTTCAAC	1557
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Qу	1738	GGCGTCCTGCTCACCCGGCCAGTCATCCTGGCTATGGACCACTGTGGGGAGCCCAGCCCT	1797
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Qу	1855	CTGCACCTGGGCGAGGAGGCGCCCTCCCACCTCTACTACTGCCAGCTGGAGGCCAGTGCC	1914
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Qу	2695	GGCCAGCCAGACGCTGGCCTCTTCACAGTGTCGGAGGCTGAGTGCTGAGGCCGGCC	752
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REFERENCE
         Venter, C.J., Adams, M.C., Li, P.W. and Myers, E.W.
 AUTHORS
 TITLE
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DEFINITION Sequence 13 from Patent WO0198354.

ACCESSION AX367094

VERSION AX367094.1 GI:18855296

KEYWORDS

SOURCE Homo sapiens (human)

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         Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
         Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
         Griffin, J.A., Kallick, D.A., Tribouley, C.M., Yue, H., Nguyen, D.B.,
 AUTHORS
         Tang, Y.T., Lal, P., Policky, J.L., Azimzai, Y., Lu, D.A., Graul, R.,
         Yao, M.G., Burford, N., Hafalia, A.J., Baughn, M.R., Bandman, O.,
         Patterson, C., Yang, J., Xu, Y., Warren, B.A., Ding, L. and
         Sanjanwala, M.S.
         Receptors
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Qу	2143	CACTTCAAGGACAGTTACCACAACCTGCGCCTATCCATCC	2202
Db	1933	CACTTCAAGGACAGTTACCACAACCTGCGCCTATCCATCC	1992
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AUTHOR TITLE	RS Engelkamp, D. Cloning of three mouse Unc5 genes and their expression patte	
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PUBME		
REFERENC	E 2	(bases 1 to 3992)

**AUTHORS** Engelkamp, D.

Direct Submission TITLE

**JOURNAL** Submitted (15-MAY-2002) Neuroanatomy, Max Planck Institute for

Brain Research, Deutschordenstrasse 46, Frankfurt 60528, GERMANY

Location/Qualifiers **FEATURES** 

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LOCUS AX268596 2697 bp DNA linear PAT 29-OCT-2001

DEFINITION Sequence 15 from Patent WO0175440.

ACCESSION AX268596

VERSION AX268596.1 GI:16541710

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SOURCE Rattus sp.

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REFERENCE AUTHOR:	S	<pre>1 (bases 1 to 2697) Leonardo, E.D., Hinck, L., Masu, M., Keino-Masu, K., Ackerman, S.L. and Tessier-Lavigne, M.</pre>					
TITLE		Vertebrate homologues of C. elegans UNC-5 are candidate netrin receptors					
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MGC:66671 IMAGE:6813463), complete cds.

ACCESSION BC058084

VERSION BC058084.1 GI:34784158

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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 3844)

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AUTHORS
            Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G.,
            Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, C.M., Schuler, G.D.,
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            Generation and initial analysis of more than 15,000 full-length
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            Email: cgapbs-r@mail.nih.gov
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            cDNA Library Preparation: M. Bento Soares, University of Iowa
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            BC Cancer Agency, Vancouver, BC, Canada
            info@bcqsc.bc.ca
            Steven Jones, Jennifer Asano, Ian Bosdet, Yaron Butterfield,
            Susanna Chan, Readman Chiu, Chris Fjell, Erin Garland, Ran Guin,
            Letticia Hsiao, Martin Krzywinski, Reta Kutsche, Oliver Lee, Soo
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 Mismatches
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           Strausberg, R.L., Feingold, E.A., Grouse, L.H., Derge, J.G.,
 AUTHORS
           Klausner, R.D., Collins, F.S., Wagner, L., Shenmen, C.M., Schuler, G.D.,
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           Generation and initial analysis of more than 15,000 full-length
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           Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
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           Strausberg, R.
 AUTHORS
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 JOURNAL
           Submitted (12-JUN-2001) National Institutes of Health, Mammalian
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Gene Collection (MGC), Cancer Genomics Office, National Cancer
            Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
            USA
            NIH-MGC Project URL: http://mgc.nci.nih.gov
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            On Dec 19, 2003 this sequence version replaced qi:14424611.
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            Akhter, N., Ayele, K., Beckstrom-Sternberg, S.M., Benjamin, B.,
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          van Criekinge, W., Roelens, I., Bogaert, T. and Verwaerde, P.
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            Ackerman, S.L., Kozak, L.P., Przyborski, S.A., Rund, L.A., Boyer, B.B.
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            The mouse rostral cerebellar malformation gene encodes an
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          Guan, W. and Condic, M.L.
 AUTHORS
          Characterization of Netrin-1, Neogenin and cUNC-5H3 expression
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## ORIGIN

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Search completed: March 6, 2005, 05:24:52 Job time: 11708.8 secs

## GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 5, 2005, 14:42:51; Search time 1401 Seconds

(without alignments)

11628.216 Million cell updates/sec

Title: US-10-624-932-1

Perfect score: 2752

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Gapop 10.0 , Gapext 1.0

Searched: 4390206 segs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

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KW
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KW
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KW
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PA
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     Alsobrook JP, Lepley DM, Gerlach VL, Macdougall JR,
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XX
     WPI; 2002-180074/23.
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     New isolated cytoplasmic, nuclear, membrane bound, or secreted
PT
     polypeptide, useful for treating cardiomyopathy, atherosclerosis,
     infections, cancer, neurodegenerative, metabolic, hematopoietic and
PT
     immune disorders.
XX
     Claim 9; Page 9-10; 213pp; English.
PS
XX
     The invention relates to an isolated cytoplasmic, nuclear, membrane
CC
CC
     bound, or secreted polypeptide (NOVX, x=1-14) their variants or mature
CC
     form. Also included are the nucleic acids encoding the NOVX proteins, a
     vector comprising the nucleic acid, a cell comprising the vector, an anti
CC
     -NOVX antibody and modulators of NOVX. NOVX, the nucleic acid and the
CC
     antibody are useful for treating or preventing a NOVX-associated
CC
CC
     disorder, where the disorder is selected from cardiomyopathy,
CC
     atherosclerosis, diabetes, a disorder related to cell signal processing
     and metabolic pathway modulation, metabolic disorders, obesity,
CC
CC
     infectious disease, anorexia, cancer-associated cachexia, cancer,
     neurodegenerative disorders, Alzheimer's disease, Parkinson's disease,
CC
```

CC immune disorders, haematopoietic disorders, and the various dyslipidaemias, metabolic disturbances associated with obesity, the CC metabolic syndrome X and wasting disorders associated with chronic CC diseases, bacterial, fungal, protozoal and viral infections, pain, CC bulimia, asthma, hypertension, urinary retention, osteoporosis, Crohn's CC disease, multiple sclerosis, Albright Hereditary Osteodystrophy, angina CC pectoris, myocardial infarction, ulcer, allergy, benign prostatic CC hypertrophy, and psychotic and neurological disorders, including anxiety, CC schizophrenia, manic depression, delirium, dementia, and dyskinesias, CC such as Huntington's disease and Gilles de la Tourette's syndrome. The CC nucleic acid is useful in gene therapy. The present sequence encodes a CC CC NOVX protein XX SQ

Sequence 2752 BP; 505 A; 937 C; 829 G; 481 T; 0 U; 0 Other;

100.0%; Score 2752; DB 6; Length 2752; Query Match Best Local Similarity 100.0%; Pred. No. 0; Matches 2752: Conservative O. Miamatahaa Indels

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Db	841	AGCCGGAGCTGCACCAACCCGGCGCCTCTCAACGGGGGCGCTTTCTGTGAGGGGCAGAAT	900
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DR
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DR
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XX
PT
    New NOVX polypeptides and nucleic acid molecules useful for preventing or
PT
     treating NOVX-associated disorders, e.g. cancer, diabetes, infection or
PT
     obesity, and in chromosome mapping, tissue typing or pharmacogenomics.
XX
PS
     Example 21; SEQ ID NO 513; 1880pp; English.
XX
CC
     The invention relates to a novel isolated polypeptide (NOVX). A
CC .
    polypeptide of the invention has cytostatic, immunomodulator,
CC
     neuroprotective, nootropic, anorectic, antidiabetic, antimicrobial, and
CC
     antilipaemic activity, and may have a use in gene therapy, and as a
CC
     vaccine. The polypeptides are encoded by NOVX polynucleotides comprising
CC
     any of the 303 fully defined nucleotide sequences given in the
CC
     specification. The polypeptide is useful in the manufacture of a
CC
    medicament for treating a syndrome associated with a human disease. The
CC
    polypeptide, polynucleotide and antibody are useful in diagnosing,
CC
     treating or preventing NOVX-associated disorders, e.g. cancer, cachexia,
CC
    Alzheimer's disease, Parkinson's disease, obesity, diabetes, infectious
CC
    diseases, metabolic syndrome X or dyslipidaemias. The nucleic acids are
CC
     further used as hybridisation probes, in chromosome mapping, tissue
CC
     typing, preventive medicine, and pharmacogenomics. The present sequence
```

encodes a NOVX polypeptide of the invention.

CC

XX

100.0%; Score 2752; DB 12; Length 2752; Query Match 100.0%; Pred. No. 0; Best Local Similarity Matches 2752; Conservative Mismatches 0; 0; 0; Indels 0; Gaps 1 CCGCGGGGCCCGCCCGGCCGCCCGCCTGCCCGCCGGGCCATGGCCGTCCGGCCC 60 Qy 1 CCGCGGGCCCCGCCCGCCCGCCCGCCGCGCGCCATGGCCGTCCGGCCC 60 Db 61 GGCCTGTGGCCAGCGCTCCTGGGCATAGTCCTCGCCGCTTTGGCTCCGCGGCTCGGGTGCC 120 Qу \_61 GGCCTGTGGCCAGCGCTCCTGGGCATAGTCCTCGCCGCTTGGCTCCGCGGCTCGGGTGCC 120 Db 121 CAGCAGAGTGCCACCGTGGCCAACCCAGTGCCTGGTGCCAACCCGGACCTGCTTCCCCAC 180 Qy 121 CAGCAGAGTGCCACCGTGGCCAACCCAGTGCCTGGTGCCAACCCGGACCTGCTTCCCCAC 180 Db 181 TTCCTGGTGGAGCCCGAGGATGTGTACATCGTCAAGAACAAGCCAGTGCTGCTTGTGTGC 240 Qу 181 TTCCTGGTGGAGCCCGAGGATGTGTACATCGTCAAGAACAAGCCAGTGCTGCTTGTGTGC 240 Db 241 AAGGCCGTGCCCGCCACGCAGATCTTCTTCAAGTGCAACGGGGAGTGGGTGCGCCAGGTG 300 Qy 241 AAGGCCGTGCCCGCCACGCAGATCTTCTTCAAGTGCAACGGGGAGTGGGTGCGCCAGGTG 300 Db 301 GACCACGTGATCGAGCGCAGCACAGACGGGAGCAGTGGGCTGCCCACCATGGAGGTCCGC 360 Qy 301 GACCACGTGATCGAGCGCAGCACAGACGGGAGCAGTGGGCTGCCCACCATGGAGGTCCGC 360 Db 361 ATTAATGTCTCAAGGCAGCAGGTCGAGAAGGTGTTCGGGCTGGAGGAATACTGGTGCCAG 420 Qу ``` 361 ATTAATGTCTCAAGGCAGCAGGTCGAGAAGGTGTTCGGGCTGGAGGAATACTGGTGCCAG 420 Db 421 TGCGTGGCATGGAGCTCCTCGGGCACCACAAGAGTCAGAAGGCCTACATCCGCATAGCC 480 Qу 421 TGCGTGGCATGGAGCTCCTCGGGCACCACCAAGAGTCAGAAGGCCTACATCCGCATAGCC 480 Db 481 AGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTGTCCCTGGAGCAGGGC 540 Qу 481 AGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTGTCCCTGGAGCAGGGC 540 Db 541 ATCGTGCTGCCTCCACCGGAGGGCATCCCTCCAGCCGAGGTGGAGTGGCTCCGG 600 Qy 541 ATCGTGCTGCCGTCCACCGGAGGGCATCCCTCCAGCCGAGGTGGAGTGGCTCCGG 600 Db 601 AACGAGGACCTGGTGGACCCGTCCCTGGACCCCAATGTATACATCACGCGGGAGCACAGC 660 Qу 601 AACGAGGACCTGGTGGACCCGTCCCTGGACCCCAATGTATACATCACGCGGGAGCACAGC 660 Db 661 CTGGTGGTGCGACAGGCCCGCCTTGCTGACACGGCCAACTACACCTGCGTGGCCAAGAAC 720 Qy 661 CTGGTGGTGCGACAGGCCCGCCTTGCTGACACGGCCAACTACACCTGCGTGGCCAAGAAC 720 Db Qу 

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	Db	781	TCGACGTGGACCGAGTGGTCCGTCTGCAGCGCCAGCTGTGGGCGCGGCTGGCAGAAACGG	840
	Qу	841	AGCCGGAGCTGCACCAACCCGGCGCCTCTCAACGGGGGCGCTTTCTGTGAGGGGCAGAAT	900
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PN
XX
PD
   25-APR-2002.
XX
PF
   15-OCT-2001; 2001WO-EP011891.
XX
PR
   16-OCT-2000; 2000US-0240061P.
XX
PA
    (FARB ) BAYER AG.
XX
PΙ
   Koehler RH;
XX
DR
   WPI; 2002-463314/49.
DR
   P-PSDB; AAU97899.
XX
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Novel human netrin binding membrane receptor polypeptide and polynucleotides for identifying modulating agents useful in treating diseases e.g. Parkinson's disease, multiple sclerosis, stroke, Alzheimer's disease.

PT XX PS

PT

PT

PT

Claim 1; Fig 1; 94pp; English.

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SO

This invention relates to the DNA and protein sequences of a novel purified human netrin binding membrane receptor, UNC5H-1. The DNA sequence of the invention is useful as a probe for detecting a nucleic acid encoding the UNC5H-1 protein in a biological sample. The sequences of the invention are useful to screen for agents which decrease the activity of the UNC5H-1 protein. The sequences are also useful for screening agents which regulate (modulate) the activity of the protein of the invention. A pharmaceutical composition containing the protein of the invention or a reagent that modulates the activity of the UNC5H-1 protein may be useful for treating a UNC5H-1 dysfunction related disease such as cancer or a central nervous system (CNS) disorders (e.g, Parkinson's disease, multiple sclerosis, stroke and Alzheimer's disease). Fusion proteins comprising the UNC5H-1 protein are useful for generating antibodies and for in various assay systems, and the protein can be used as a bait protein in a two-hybrid assay or three-hybrid assay. The method of the invention is useful for detecting a coding sequence for the UNC5H-1 protein. The present sequence represents a DNA sequence encoding the human netrin binding membrane receptor UNC5H-1 protein of the invention

Sequence 2697 BP; 503 A; 906 C; 807 G; 481 T; 0 U; 0 Other;

Query Match 97.7%; Score 2687.4; DB 6; Length 2697; Best Local Similarity 99.8%; Pred. No. 0; Matches 2691; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

46 ATGGCCGTCCGGCCCGGCCTGTGGCCAGCGCTCCTGGGCATAGTCCTCGCCGCTTGGCTC 105 Qу 1 ATGGCCGTCCGGCCCGGCCTGTGGCCAGCGCTCCTGGGCATAGTCCTCGCCGCTTGGCTC 60 Db 106 CGCGGCTCGGGTGCCCAGCAGAGTGCCACCGTGGCCAACCCAGTGCCTGGTGCCAACCCG 165 QУ 61 CGCGGCTCGGGTGCCCAGCAGAGTGCCACCGTGGCCAACCCAGTGCCTGGTGCCAACCCG 120 Db 166 GACCTGCTTCCCCACTTCCTGGTGGAGCCCGAGGATGTGTACATCGTCAAGAACAAGCCA 225 Qу 121 GACCTGCTTCCCCACTTCCTGGTGGAGCCCGAGGATGTGTACATCGTCAAGAACAAGCCA 180 Db 226 GTGCTGCTGTGCAAGGCCGTGCCCGCCACGCAGATCTTCTTCAAGTGCAACGGGGAG 285 Qy 181 GTGCTGCTTGTGCAAGGCCGTGCCCGCCACGCAGATCTTCTTCAAGTGCAACGGGGAG 240 Dh 286 TGGGTGCGCCAGGTGGACCACGTGATCGAGCGCAGCACAGACGGGAGCAGTGGGCTGCCC 345 Qу 241 TGGGTGCGCCAGGTGGACCACGTGATCGAGCGCAGCAGACGGGAGCAGTGGGCTGCCC 300 Db 346 ACCATGGAGGTCCGCATTAATGTCTCAAGGCAGCAGGTCGAGAAGGTGTTCGGGCTGGAG 405 Qу 301 ACCATGGAGGTCCGCATTAATGTCTCAAGGCAGCAGGTCGAGAAGGTGTTCGGGCTGGAG 360 Db 406 GAATACTGGTGCCAGTGCGTGGCATGGAGCTCCTCGGGCACCACCAAGAGTCAGAAGGCC 465 Qy

Db	361	GAATACTGGTGCCAGTGCGTGGCATGGAGCTCCTCGGGCACCAAGAGTCAGAAGGCC	420
Qy	466	TACATCCGCATAGCCAGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTG	525
Db	421	TACATCCGCATAGCCTATTTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTG	480
Qy	526	TCCCTGGAGCAGGGCATCGTGCTGCCCTGCCGTCCACCGGAGGGCATCCCTCCAGCCGAG	585
Db	481	TCCCTGGAGCAGGGCATCGTGCCGTGCCGTCCACCGGAGGGCATCCCTCCAGCCGAG	540
Qу	586	GTGGAGTGGCTCCGGAACGAGGACCTGGTGGACCCGTCCCTGGACCCCAATGTATACATC	645
Db	541	GTGGAGTGGCTCCGGAACGAGGACCTGGTGGACCCGTCCCTGGACCCCAATGTATACATC	600
Qу	646	ACGCGGGAGCACAGCCTGGTGGTGCGACAGGCCCGCCTTGCTGACACGGCCAACTACACC	705
Db	601		660
Qу	706	TGCGTGGCCAAGAACATCGTGGCACGTCGCCGCAGCGCCTCCGCTGTCATCGTCTAC	765
Db	661	TGCGTGGCCAAGAACATCGTGGCACGTCGCCGCAGCGCCTCCGCTGCTGTCATCGTCTAC	720
Qγ	766	GTGAACGGTGGGTCGACGTGGACCGAGTGGTCCGTCTGCAGCGCCAGCTGTGGGCCC	825
Db	721	GTGAACGGTGGGTCGACGTGGACCGAGTGGTCCGTCTGCAGCGCCAGCTGTGGGCGC	780
Qy	826	GGCTGGCAGAACGGAGCCGGAGCTGCACCAACCCGGCGCCTCTCAACGGGGGCGCTTTC	885
Db	781	GGCTGGCAGAAACGGAGCCGGAGCTGCACCAACCCGGCGCCTCTCAACGGGGGCGCTTTC	840
Qу	886	TGTGAGGGCAGAATGTCCAGAAAACAGCCTGCGCCACCCTGTGCCCAGTAGACGGCAGC	945
Db	841	TGTGAGGGCAGAATGTCCAGAAAACAGCCTGCGCCACCCTGTGCCCAGTGGACGGCAGC	900
Qу	946	TGGAGCCCGTGGAGCAAGTGGTCGGCCTGTGGGCTGGACTGCACCCACTGGCGGAGCCGT	1005
Db	901	TGGAGCCCGTGGAGCAAGTGGTCGGCCTGTGGGCTGGACTGCACCCACTGGCGGAGCCGT	960
Qу	1006	GAGTGCTCTGACCCAGCACCCCGCAACGGAGGGAGGAGTGCCAGGGCACTGACCTGGAC	1065
Db	961	GAGTGCTCTGACCCAGCACCCCGCAACGGAGGGAGGGAGTGCCAGGGCACTGACCTGGAC	1020
Qу	1066	ACCCGCAACTGTACCAGTGACCTCTGTGTACACAGTGCTTCTGGCCCTGAGGACGTGGCC	1125
Db	1021	ACCCGCAACTGTACCAGTGACCTCTGTGTACACACTGCTTCTGGCCCTGAGGACGTGGCC	1080
Qу	1126	CTCTATGTGGGCCTCATCGCCGTGGCCGTCTGCCTGCTGCTGCTGCTGCTGCTCATC	1185
Db	1081	CTCTATGTGGGCCTCATCGCCGTGGCCGTCTGCCTGCTGCTGCTGCTCATC	1140
Qу	1186	CTCGTTTATTGCCGGAAGAAGGAGGGGCTGGACTCAGATGTGGCTGACTCGTCCATTCTC	1245
Db	1141	CTCGTTTATTGCCGGAAGAAGGAGGGGCTGGACTCAGATGTGGCTGACTCGTCCATTCTC	1200
Qу	1246	ACCTCAGGCTTCCAGCCCGTCAGCATCAAGCCCAGCAAAGCAGACAACCCCCATCTGCTC	1305

Dp.	1201	${\tt ACCTCAGGCTTCCAGCCCGTCAGCATCAAGCCCAGCAAAGCAGACAACCCCCATCTGCTC}$	1260
Qу	1306	ACCATCCAGCCGGACCTCAGCACCACCACCACCACCAGGGCAGTCTCTGTCCCCGG	1365
Db	1261	ACCATCCAGCCGGACCTCAGCACCACCACCACCAGGGCAGTCTCTGTCCCCGG	1320
Qy	1366	CAGGATGGGCCCAGCCCCAAGTTCCAGCTCACCAATGGGCACCTGCTCAGCCCCCTGGGT	1425
Db	1321	CAGGATGGGCCCAGCCCCAAGTTCCAGCTCACCAATGGGCACCTGCTCAGCCCCCTGGGT	1380
Qy	1426	GGCGGCCGCCACACTGCACCACAGCTCTCCCACCTCTGAGGCCGAGGAGTTCGTCTCC	1485
Db	1381	GGCGGCCGCCACACTGCACCACACTCTCCCACCTCTGAGGCCGAGGAGTTCGTCTCC	1440
Qy	1486	CGCCTCTCCACCCAGAACTACTTCCGCTCCCTGCCCCGAGGCACCAGCAACATGACCTAT	1545
Db	1441	CGCCTCTCCACCCAGAACTACTTCCGCTCCCTGCCCCGAGGCACCAGCAACATGACCTAT	1500
Qy	1546	GGGACCTTCAACTTCCTCGGGGGCCGGCTGATGATCCCTAATACAGGTATCAGCCTCCTC	1605
Db	1501	GGGACCTTCAACTTCCTCGGGGGCCGGCTGATGATCCCTAATACAGGAATCAGCCTCCTC	1560
Qy	1606	ATCCCCCCAGATGCCATACCCCGAGGGAAGATCTATGAGATCTACCTCACGCTGCACAAG	1665
Db	1561	ATCCCCCCAGATGCCATACCCCGAGGGAAGATCTATGAGATCTACCTCACGCTGCACAAG	1620
Qy	1666	CCGGAAGACGTGAGGTTGCCCCTAGCTGGCTGTCAGACCCTGCTGAGTCCCATCGTTAGC	1725
Db	1621	CCGGAAGACGTGAGGTTGCCCCTAGCTGGCTGTCAGACCCTGCTGAGTCCCATCGTTAGC	1680
Qy	1726	TGTGGACCCCTGGCGTCCTGCTCACCCGGCCAGTCATCCTGGCTATGGACCACTGTGGG	1785
Db	1681	TGTGGACCCCTGGCGTCCTGCTCACCCGGCCAGTCATCCTGGCTATGGACCACTGTGGG	1740
Qу	1786	GAGCCCAGCCCTGACAGCTGGAGCCTGCGCCTCAAAAAGCAGTCGTGCGAGGGCAGCTGG	1845
Db	1741	GAGCCCAGCCTGACAGCTGGAGCCTGCGCCTCAAAAAGCAGTCGTGCGAGGGCAGCTGG	1800
Qу	1846	GAGGATGTGCTGCACCTGGGCGAGGAGGCGCCCTCCCACCTCTACTACTGCCAGCTGGAG	1905
Db	1801	GAGGATGTGCTGCACCTGGGCGAGGAGGCGCCCTCCCACCTCTACTACTGCCAGCTGGAG	1860
Qу	1906	GCCAGTGCCTGCTACGTCTTCACCGAGCAGCTGGGCCGCTTTGCCCTGGTGGGAGAGGCC	1965
Db	1861	GCCAGTGCCTGCTACGTCTTCACCGAGCAGCTGGGCCGCTTTGCCCTGGTGGGAGAGGCC	1920
Qy	1966	CTCAGCGTGGCTGCCCCAAGCGCCTCAAGCTGCTTCTGTTTGCGCCGGTGGCCTGCACC	2025
Db	1921	CTCAGCGTGGCTGCCCAAGCGCCTCAAGCTGCTTCTGTTTGCGCCGGTGGCCTGCACC	1980
Qy	2026	TCCCTCGAGTACAACATCCGGGTCTACTGCCTGCATGACACCCACGATGCACTCAAGGAG	2085
Db	1981	TCCCTCGAGTACAACATCCGGGTCTACTGCCTGCATGACACCCACGATGCACTCAAGGAG	2040
Qу	2086	GTGGTGCAGCTGGAGAAGCAGCTGGGGGGACAGCTGATCCAGGAGCCACGGGTCCTGCAC	2145
Db ·	2041	GTGGTGCAGCTGGAGAAGCAGCTGGGGGGACAGCTGATCCAGGAGCCACGGGTCCTGCAC	2100

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2101 TTCAAGGACAGTTACCACAACCTGCGCCTATCCATCCACGATGTGCCCAGCTCCCTGTGG 2160
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      2206 AAGAGTAAGCTCCTTGTCAGCTACCAGGAGATCCCCTTTTATCACATCTGGAATGGCACG 2265
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         2161 AAGAGTAAGCTCCTTGTCAGCTACCAGGAGATCCCCTTTTATCACATCTGGAATGGCACG 2220
Db
      2266 CAGCGGTACTTGCACTGCACCTTGACCCTGGAGCGTGTCAGCCCCAGCACTAGTGACCTG 2325
Qу
         2221 CAGCGGTACTTGCACTGCACCTTCACCCTGGAGCGTGTCAGCCCCAGCACTAGTGACCTG 2280
Db
      2326 GCCTGCAAGCTGTGGGTGGGCAGGTGGAGGGCGACGGGCAGAGCTTCAGCATCAACTTC 2385
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         2281 GCCTGCAAGCTGTGGGTGGGAGGTGGAGGGCGACGGGCAGAGCTTCAGCATCAACTTC 2340
Db
      2386 AACATCACCAAGGACACAAGGTTTGCTGAGCTGCTGGCTCTGGAGAGTGAAGCGGGGGTC 2445
Qу
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      2446 CCAGCCTGGTGGGCCCCAGTGCCTTCAAGATCCCCTTCCTCATTCGGCAGAAGATAATT 2505
Qу
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         2461 TCCAGCCTGGACCCACCCTGTAGGCGGGGTGCCGACTGGCGGACTCTGGCCCAGAAACTC 2520
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         Db
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RESULT 4
ABK49422
ID
   ABK49422 standard; DNA; 2881 BP.
XX
AC
   ABK49422;
XX
DT
   15-JUL-2002
            (first entry)
XX
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   DNA encoding human UNC5-like protein NOV1.
XX
KW
   Human; NOVX polypeptide; cardiomyopathy; atherosclerosis; cancer;
KW
   cell signal processing; metabolic pathway modulation; cancerous tissue;
   antibody; diabetes; transgenic animal; UNC5-like protein; NOV1;
KW
KW
   chromosome 13; gene; ds.
XX
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2146 TTCAAGGACAGTTACCACAACCTGCGCCTATCCATCCACGATGTGCCCAGCTCCCTGTGG 2205

Qу

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Homo sapiens.
OS
XX
                   Location/Oualifiers
FH
    Key
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    CDS
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                   /product= "Human UNC5-like protein NOV1"
XX
PN
    WO200229038-A2.
XX
PD
    11-APR-2002.
XX
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    04-OCT-2001; 2001WO-US031377.
XX
    04-OCT-2000; 2000US-0237862P.
PR
XX
PA
     (CURA-) CURAGEN CORP.
XX
PΙ
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                             Shimkets RA;
XX
DR
    WPI; 2002-340104/37.
DR
    P-PSDB; AAU79939.
XX
PΤ
    Novel isolated NOVX polypeptide, and encoded polynucleotide, useful for
PT
    treating cardiomyopathy, artherosclerosis, and cancer.
XX
PS
    Claim 8; Page 7-8; 180pp; English.
XX
CC
    The present invention relates to a new NOVX polypeptide having a 900
     (NOV1), 4349 (NOV2), 940 (NOV3), 798 (NOV4), 865 (NOV5), or 331 (NOV6)
CC
    residue amino acid sequence, as given in the specification. The novel
CC
CC
    polypeptide, and its encoding polynucleotide, are used to treat
CC
    cardiomyopathy, atherosclerosis, cancer or a disease related to cell
    signal processing and metabolic pathway modulation, in a human. Detecting
CC
CC
    the polypeptide or polynucleotide is useful for identifying cancerous
    tissue. The antibody can be used to treat diabetes or cancer. The host
CC
    cells can be used to produce non-human transgenic animals useful in drug
CC
    screening. The present nucleic acid sequence is that of the human UNC5-
CC
    like NOV1 gene located on chromosome 13. This sequence encodes the human
CC
    UNC5-like protein NOV1 of the invention
CC
XX
    Sequence 2881 BP; 526 A; 985 C; 868 G; 502 T; 0 U; 0 Other;
SO
                               Score 2676.4; DB 6;
                        97.3%;
                                                   Length 2881;
  Query Match
                        98.9%;
                               Pred. No. 0;
  Best Local Similarity
 Matches 2728; Conservative
                              0; Mismatches
                                            21;
                                                  Indels
                                                                       3;
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          42 CCGCGGGCCCCGCCCGCCCGCCCGCCCGCCGGCCATGGCCGTCCGGCCC 101
          61 GGCCTGTGGCCAGCGCTCCTGGGCATAGTCCTCGCCGCTTGGCTCCGCGGCTCGGGTGCC 120
Qу
             102 GGCCTGTGGCCAGCGCTCCTGGGCATAGTCCTCGCCGCTTGGCTCCGCGGCTCGGGTGCC 161
Db
         121 CAGCAGAGTGCCACCGTGGCCAACCCAGTGCCTGGTGCCAACCCGGACCTGCTTCCCCAC 180
Qу
             Db
         162 CAGCAGAGTGCCACCGTGGCCAACCCAGTGCCTGGTGCCAACCCGGACCTGCTTCCCCAC 221
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Qy	181	TTCCTGGTGGAGCCCGAGGATGTGTACATCGTCAAGAACAAGCCAGTGCTGCTTGTGTGC	240
Db	222	${\tt TTCCTGGTGGAGCCCGAGGATGTGTACATCGTCAAGAACAAGCCAGTGCTGCTTGTGTGCCTGCTGTGTGTG$	281
Qy	241	AAGGCCGTGCCCGCCACGCAGATCTTCTTCAAGTGCAACGGGGAGTGGGTGCGCCAGGTG	300
Db	282	AAGGCCGTGCCCGCCACGCAGATCTTCTTCAAGTGCAACGGGGAGTGGGTGCGCCAGGTG	341
Qу	301	GACCACGTGATCGAGCGCAGCACAGACGGGAGCAGTGGGCTGCCCACCATGGAGGTCCGC	360
Db	342	GACCACGTGATCGAGCGCAGCAGACGGGAGCAGTGGTGAGCCGACCATGGAGGTCCGC	401
Qу	361	ATTAATGTCTCAAGGCAGCAGGTCGAGAAGGTGTTCGGGCTGGAGGAATACTGGTGCCAG	420
Db	402	ATTAATGTCTCAAGGCAGCAGGTCGAGAAGGTGTTCGGGCTGGAGGAATACTGGTGCCAG	461
Qy .	421	TGCGTGGCATGGAGCTCCTCGGGCACCACCAAGAGTCAGAAGGCCTACATCCGCATAGCC	480
Db	462	TGCGTGGCATGGAGCTCCTCGGGCACCACCAAGAGTCAGAAGGCCTACATCCGCATAGCC	521
Qy	481	AGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTGTCCCTGGAGCAGGGC	540
ĎЬ	522	AGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTGTCCCTGGAGCAGGGC	581
Qy	541	ATCGTGCTGCCCTCCACCGGAGGGCATCCCTCCAGCCGAGGTGGAGTGGCTCCGG	600
Db	582	ATCGTGCTGCCCTCCACCGGAGGGCATCCCTCCAGCCGAGGTGGAGTGGCTCCGG	641
Qy	601	AACGAGGACCTGGTGGACCCGTCCCTGGACCCCAATGTATACATCACGCGGGAGCACAGC	660
Db	642	AACGAGGACCTGGTGGACCCGTCCCTGGACCCCAATGTATACATCACGCGGGAGCACAGC	701
Qy	661	CTGGTGGTGCGACAGGCCCGCCTTGCTGACACGGCCAACTACACCTGCGTGGCCAAGAAC	720
Db	702	CTGGTGGTGCGACAGGCCCGCCTTGCTGACACGGCCAACTACACCTGCGTGGCCAAGAAC	761
Qу	721	ATCGTGGCACGTCGCCGCAGCGCCTCCGCTGCTGTCATCGTCAACGTGAACGGTGGGTG	780
Db	762	ATCGTGGCACGTCGCCGCAGCGCCTCCGCTGTCATCGTCTACGTGAACGGTGGGTG	821
Qy	781	TCGACGTGGACCGAGTGGTCCGTCTGCAGCGCCAGCTGTGGGCGCGGCTGGCAGAAACGG	840
Db	822	TCGACGTGGACCGAGTGGTCCGTCTGCAGCGCCAGCTGTGGGCGCGGCTGGCAGAAACGG	881
Qy	841	AGCCGGAGCTGCACCAACCCGGCGCCTCTCAACGGGGGCGCTTTCTGTGAGGGGCAGAAT	900
Db	882	AGCCGGAGCTGCACCAACCCGGCGCCTCTCAACGGGGGCGCTTTCTGTGAGGGGCAGAAT	941
Qy	901	GTCCAGAAAACAGCCTGCGCCACCCTGTGCCCAGTAGACGGCAGCTGGAGCCCGTGG	957
Db	942		1001
Qу	958	AGCAAGTGGTCGGCCTGTGGGCTGGACTGCACCCACTGGCGGAGCCGTGAGTGCTCTGAC	1017
Db :	1002	AGCAAGTGGTCGGCCTGTGGGCTGGACTGCACCCACTGGCGGAGCCGTGAGTGCTCTGAC	1061

Qу	1018	CCAGCACCCGCAACGGAGGGAGGAGTGCCAGGGCACTGACCTGGACACCCGCAACTGT	1077
Db	1062		1121
Qy	1078	ACCAGTGACCTCTGTGTACACAGTGCTTCTGGCCCTGAGGACGTGGCCCTCTATGTGGGC	1137
Db	1122	ACCAGTGACCTCTGTGTACACAGTGCTTCTGGCCCTGAGGACGTGGCCCTCTATGTGGGC	1181
Qу	1138	CTCATCGCCGTGGCCGTCTGCCTGCTGCTGCTGCTGCTCATCCTCGTTTATTGC	1197
Db	1182		1241
Qу	1198	CGGAAGAAGGAGGGCTGGACTCAGATGTGGCTGACTCGTCCATTCTCACCTCAGGCTTC	1257
Db	1242		1301
Qy	1258	CAGCCCGTCAGCATCAAGCCCAGCAAAGCAGACAACCCCCATCTGCTCACCATCCAGCCG	1317
Db	1302	CAGCCCGTCAGCATCAAGCCCAGCAAAGCAGACAACCCCCATCTGCTCACCATCCAGCCG	1361
Qy	1318	GACCTCAGCACCACCACCACCTACCAGGGCAGTCTCTGTCCCCGGCAGGATGGGCCC	1377
Db	1362	GACCTCAGCACCACCACCACCTACCAGGGCAGTCTCTGTCCCCGGCAGGATGGGCCC	1418
Qy	1378	AGCCCCAAGTTCCAGCTCACCAATGGGCACCTGCTCAGCCCCCTGGGTGGCGGCCGCCAC	1437
Db	1419	AGCCCCAAGTTCCAGCTCACCAATGGGCACCTGCTCAGCCCCCTGGGTGGCGGCCGCCAC	1478
Qу	1438	ACACTGCACCACGCTCTCCACCTCTGAGGCCGAGGAGTTCGTCTCCCGCCTCTCCACC	
Db	1479	ACACTGCACCACAGCTCTCCACCTCTGAGGCCGAGGAGTTCGTCTCCCGCCTCTCCACC	
Qу	1498	CAGAACTACTTCCGCTCCCTGCCCCGAGGCACCAGCAACATGACCTATGGGACCTTCAAC	1557
Db	1539	CAGAACTACTTCCGCTCCCTGCCCCGAGGCACCAGCAACATGACCTATGGGACCTTCAAC	1598
Qy.	1558	TTCCTCGGGGGCCGGCTGATGATCCCTAATACAGGTATCAGCCTCCTCATCCCCCAGAT	1617
Db	1599	TTCCTCGGGGGCCGGCTGATGATCCCTAATACAGGTATCAGCCTCCTCATCCCCCAGAT	1658
Qу	1618	GCCATACCCCGAGGGAAGATCTATGAGATCTACCTCACGCTGCACAAGCCGGAAGACGTG	1677
Db	1659	GCCATACCCCGAGGGAAGATCTATGAGATCTACCTCACGCTGCACAAGCCGGAAGACGTG	1718
Qу	1678	AGGTTGCCCCTAGCTGGCTGTCAGACCCTGCTGAGTCCCATCGTTAGCTGTGGACCCCCT	1737
Db	1719	AGGTTGCCCCTAGCTGGCTGTCAGACCCTGCTGAGTCCCATCGTTAGCTGTGGACCCCCT	1778
Qу	1738	GGCGTCCTGCTCACCCGGCCAGTCATCCTGGCTATGGACCACTGTGGGGAGCCCAGCCCT	1797
Db	1779	GGCGTCCTGCTCACCCGGCCAGTCATCCTGGCTATGGACCACTGTGGGGAGCCCAGCCCT	1838
Qy	1798	GACAGCTGGAGCCTGCGCCTCAAAAAGCAGTCGTGCGAGGGCAGCTGGGAGGATGTG	1854
Db	1839	GACAGCTGGAGCCTCCAAAAAGCAGTCGTGCGAGGGCAGCTGGGAGCAGGATGTG	1898
Ov	1855	CTGCACCTGGGCGAGGAGGCGCCCTCCCACCTCTACTACTGCCAGCTGGAGGCCAGTGCC	1914

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Db	1899		19,58
Qу	1915	TGCTACGTCTTCACCGAGCAGCTGGGCCGCTTTGCCCTGGTGGGAGAGGCCCTCAGCGTG	1974
Db	1959	TGCTACGTCTTCACCGAGCAGCTGGGCCGCTTTGCCCTGGTGGGAGAGGCCCTCAGCGTG	2018
Qу	1975	GCTGCCGCCAAGCGCCTCAAGCTGCTTCTGTTTGCGCCGGTGGCCTGCACCTCCCTC	2034
Db	2019		2078
Qу	2035	TACAACATCCGGGTCTACTGCCTGCATGACACCCACGATGCACTCAAGGAGGTGGTGCAG	2094
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Qy	2095	CTGGAGAAGCAGCTGGGGGGACAGCTGATCCAGGAGCCACGGGTCCTGCACTTCAAGGAC	2154
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Qу	2155	AGTTACCACAACCTGCGCCTATCCATCCACGATGTGCCCAGCTCCCTGTGGAAGAGTAAG	2214
Db	2199	AGTTACCACAACCTGCGCCTATCCATCCACGATGTGCCCAGCTCCCTGTGGAAGAGTAAG	2258
Qу	2215	CTCCTTGTCAGCTACCAGGAGATCCCCTTTTATCACATCTGGAATGGCACGCAGCGGTAC	2274
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Qу	2275	TTGCACTGCACCTTCACCCTGGAGCGTGTCAGCCCCAGCACTAGTGACCTGGCCTGCAAG	2334
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Qу	2455	GTGGGCCCCAGTGCCTTCAAGATCCCCTTCCTCATTCGGCAGAAGATAATTTCCAGCCTG	2514
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Qу	2515	GACCCACCCTGTAGGCGGGGTGCCGACTGGCGGACTCTGGCCCAGAAACTCCACCTGGAC	2574
Db	2559	GACCCACCCTGTAGGCGGGTGCCGACTGGCGGACTCTGGCCCAGAAACTCCACCTGGAC	2618
Qу	2575	AGCCATCTCAGCTTCTTTGCCTCCAAGCCCAGCCCCACAGCCATGATCCTCAACCTGTGG	2634
Db	2619	AGCCATCTCAGCTTCTTTGCCTCCAAGCCCAGCCCCACAGCCATGATCCTCAACCTGTGG	2678
Qу	2635	GAGGCGCGCACTTCCCCAACGGCAACCTCAGCCAGCTGGCTG	2694
Db	2679	GAGGCGCGCACTTCCCCAACGGCAACCTCAGCCAGCTGGCTG	2738
Qу	2695	GGCCAGCCAGACGCTGGCCTCTTCACAGTGTCGGAGGCTGAGTGCTGAGGCCGGCC	752

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RESULT 5
ADH71609
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ID .
XX
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XX
DT
     25-MAR-2004
                  (first entry)
XX
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XX
KW
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KW
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KW
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KW
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XX
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    New NOVX polypeptides and nucleic acid molecules useful for preventing or
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    obesity, and in chromosome mapping, tissue typing or pharmacogenomics.
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    The invention relates to a novel isolated polypeptide (NOVX). A
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    polypeptide of the invention has cytostatic, immunomodulator,
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    neuroprotective, nootropic, anorectic, antidiabetic, antimicrobial, and
CC
    antilipaemic activity, and may have a use in gene therapy, and as a
CC
CC
    vaccine. The polypeptides are encoded by NOVX polynucleotides comprising
    any of the 303 fully defined nucleotide sequences given in the
CC
    specification. The polypeptide is useful in the manufacture of a
CC
    medicament for treating a syndrome associated with a human disease. The
CC
    polypeptide, polynucleotide and antibody are useful in diagnosing,
CC
    treating or preventing NOVX-associated disorders, e.g. cancer, cachexia,
CC
    Alzheimer's disease, Parkinson's disease, obesity, diabetes, infectious
CC
    diseases, metabolic syndrome X or dyslipidaemias. The nucleic acids are
CC
    further used as hybridisation probes, in chromosome mapping, tissue
CC
    typing, preventive medicine, and pharmacogenomics. The present sequence
CC
    encodes a NOVX polypeptide of the invention.
CC
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                                                         9; Gaps
                                                                   3;
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Qу
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            222 TTCCTGGTGGAGCCCGAGGATGTGTACATCGTCAAGAACAAGCCAGTGCTGCTTGTGTGC 281
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            Db
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Qу	421	TGCGTGGCATGGAGCTCCTCGGGCACCACCAAGAGTCAGAAGGCCTACATCCGCATAGCC	480
Db	462	TGCGTGGCATGGAGCTCCTCGGGCACCACCAAGAGTCAGAAGGCCTACATCCGCATAGCC	521
Qу	481	AGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTGTCCCTGGAGCAGGGC	540
Db	522	AGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTGTCCCTGGAGCAGGGC	581
Qу	541	ATCGTGCTGCCGTCCACCGGAGGGCATCCCTCCAGCCGAGGTGGAGTGGCTCCGG	600
Db	582	ATCGTGCTGCCGTCCACCGGAGGGCATCCCTCCAGCCGAGGTGGAGTGGCTCCGG	641
QУ	601	AACGAGGACCTGGTGGACCCGTCCCTGGACCCCAATGTATACATCACGCGGGAGCACAGC	660
Db	642	AACGAGGACCTGGTGGACCCCTGGACCCCAATGTATACATCACGCGGGAGCACAGC	701
Qу	661	CTGGTGGTGCGACAGGCCCGCCTTGCTGACACGGCCAACTACACCTGCGTGGCCAAGAAC	720
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Qy	721	ATCGTGGCACGTCGCCGCAGCGCCTCCGCTGTCATCGTCTACGTGAACGGTGGGTG	780
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Qу	781	TCGACGTGGACCGAGTGGTCCGTCTGCAGCGCCAGCTGTGGGCGCGGCTGGCAGAAACGG	840
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Qу	841	AGCCGGAGCTGCACCAACCCGGCGCCTCTCAACGGGGGCGCTTTCTGTGAGGGGCAGAAT	900
Db	882	AGCCGGAGCTGCACCAACCCGGCGCCTCTCAACGGGGGCGCTTTCTGTGAGGGGCAGA	941
Qу	901	GTCCAGAAAACAGCCTGCGCCACCCTGTGCCCAGTAGACGGCAGCTGGAGCCCGTGG	957
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Qу	958	AGCAAGTGGTCGGCCTGTGGGCTGGACTGCACCCACTGGCGGAGCCGTGAGTGCTCTGAC	1017
Db	1002	AGCAAGTGGTCGGCCTGTGGGCTGGACTGCACCCACTGGCGGAGCCGTGAGTGCTCTGAC	1061
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Qу	1855	CTGCACCTGGGCGAGGAGGCGCCCTCCCACCTCTACTACTGCCAGCTGGAGGCCAGTGCC	1914
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Qу	1915	TGCTACGTCTTCACCGAGCAGCTGGGCCGCTTTGCCCTGGTGGGAGAGGCCCTCAGCGTG	1974
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Qу	1975	GCTGCCGCCAAGCGCCTCAAGCTGCTTCTGTTTGCGCCGGTGGCCTGCACCTCCCTC	2034
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25-MAR-2004

(first entry)

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     treating NOVX-associated disorders, e.g. cancer, diabetes, infection or
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     obesity, and in chromosome mapping, tissue typing or pharmacogenomics.
XX
PS
     Example 21; SEQ ID NO 545; 1880pp; English.
XX
CC
     The invention relates to a novel isolated polypeptide (NOVX). A
CC
     polypeptide of the invention has cytostatic, immunomodulator,
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neuroprotective, nootropic, anorectic, antidiabetic, antimicrobial, and CC antilipaemic activity, and may have a use in gene therapy, and as a CC vaccine. The polypeptides are encoded by NOVX polynucleotides comprising CC any of the 303 fully defined nucleotide sequences given in the CC specification. The polypeptide is useful in the manufacture of a CC medicament for treating a syndrome associated with a human disease. The CC polypeptide, polynucleotide and antibody are useful in diagnosing, CC treating or preventing NOVX-associated disorders, e.g. cancer, cachexia, CC CC Alzheimer's disease, Parkinson's disease, obesity, diabetes, infectious diseases, metabolic syndrome X or dyslipidaemias. The nucleic acids are CC further used as hybridisation probes, in chromosome mapping, tissue CC CC typing, preventive medicine, and pharmacogenomics. The present sequence CC encodes a NOVX polypeptide of the invention. XX

SQ Sequence 2881 BP; 526 A; 986 C; 868 G; 501 T; 0 U; 0 Other;

Query Match 97.2%; Score 2674.8; DB 12; Length 2881; Best Local Similarity 98.9%; Pred. No. 0; Matches 2727; Conservative 0; Mismatches 22; Indels 9; Gaps 3;

	Matches	212	7; Conservative 0; Mismatches 22; Indels 9; Gaps	3
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D	b	102		161
Q	У	121	CAGCAGAGTGCCACCGTGGCCAACCCAGTGCCTGGTGCCAACCCGGACCTGCTTCCCCAC	180
D	b	162		221
Q	У	181	TTCCTGGTGGAGCCCGAGGATGTGTACATCGTCAAGAACAAGCCAGTGCTGCTTGTGTGC	240
D	b	222	TTCCTGGTGGAGCCCGAGGATGTGTACATCGTCAAGAACAAGCCAGTGCTGCTTGTGTGC	281
Q	У	241	AAGGCCGTGCCCGCCACGCAGATCTTCTTCAAGTGCAACGGGGAGTGGGTGCGCCAGGTG	300
D	b	282	AAGGCCGTGCCCGCCACGCAGATCTTCTTCAAGTGCAACGGGGAGTGGGTGCGCCAGGTG	341
Q	У	301	GACCACGTGATCGAGCGCAGCACAGACGGGAGCAGTGGGCTGCCCACCATGGAGGTCCGC	360
D	b	342		401
Q	у	361	ATTAATGTCTCAAGGCAGCAGGTCGAGAAGGTGTTCGGGCTGGAGGAATACTGGTGCCAG	420
D	b	402		461
Q	У	421	TGCGTGGCATGGAGCTCCTCGGGCACCACCAAGAGTCAGAAGGCCTACATCCGCATAGCC	480
D	b	462	TGCGTGGCATGGAGCTCCTCGGGCACCACCAAGAGTCAGAAGGCCTACATCCGCATAGCC	521
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Db	582	ATCGTGCTGCCCTGCCGTCCACCGGAGGCATCCCTCCAGCCGAGGTGGAGTGGCTCCGG	641
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Db	762	ATCGTGGCACGTCGCCGCAGCGCTCCGCTGTCATCGTCTACGTGAACGGTGGGTG	821
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Db	882	AGCCGGAGCTGCACCAACCCGGCGCCTCTCAACGGGGGCGCTTTCTGTGAGGGGCAGAAT	941
Qу	901	GTCCAGAAAACAGCCTGCGCCACCCTGTGCCCAGTAGACGGCAGCTGGAGCCCGTGG	957
Db	942	GTCCATGACCGCACCGTCTCCTCTCTGCTTGTCTCTGTGGACGGCAGCTGGAGCCCGTGG	1001
Qу	958	AGCAAGTGGTCGGCCTGTGGGCTGGACTGCACCCACTGGCGGAGCCGTGAGTGCTCTGAC	1017
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Qу	1018	CCAGCACCCGCAACGGAGGGGAGGGGGGGGGGGGGCACTGACCTGGACACCCGCAACTGT	1077
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Db	1182	CTCATCGCCGTGGCCGTCTGCCTGGTCCTGCTGCTGCTTGTCCTCATCCTCGTTTATTGC	1241
Qу	1198	CGGAAGAAGGAGGGGCTGGACTCAGATGTGGCTGACTCGTCCATTCTCACCTCAGGCTTC	1257
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Qу	1258	CAGCCCGTCAGCATCAAGCCCAGCAAAGCAGACAACCCCCATCTGCTCACCATCCAGCCG	1317
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Qу	1318	GACCTCAGCACCACCACCACCTACCAGGGCAGTCTCTGTCCCCGGCAGGATGGGCCC	1377
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Db	1419	AGCCCCAAGTTCCAGCTCACCAATGGGCACCTGCTCAGCCCCCTGGGTGGCGGCCGCCAC	1478
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Qу	1618	GCCATACCCCGAGGGAAGATCTATGAGATCTACCTCACGCTGCACAAGCCGGAAGACGTG	1677
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Qу		CTCCTTGTCAGCTACCAGGAGATCCCCTTTTATCACATCTGGAATGGCACGCAGCGGTAC	
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XX
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     New NOVX polypeptides and nucleic acid molecules useful for preventing or
PT
     treating NOVX-associated disorders, e.g. cancer, diabetes, infection or
PT
     obesity, and in chromosome mapping, tissue typing or pharmacogenomics.
XX
PS
     Example 21; SEQ ID NO 531; 1880pp; English.
XX
     The invention relates to a novel isolated polypeptide (NOVX). A
CC
CC
     polypeptide of the invention has cytostatic, immunomodulator,
     neuroprotective, nootropic, anorectic, antidiabetic, antimicrobial, and
CC
CC
     antilipaemic activity, and may have a use in gene therapy, and as a
     vaccine. The polypeptides are encoded by NOVX polynucleotides comprising
CC
CC
     any of the 303 fully defined nucleotide sequences given in the
     specification. The polypeptide is useful in the manufacture of a
CC
CC
     medicament for treating a syndrome associated with a human disease. The
CC
     polypeptide, polynucleotide and antibody are useful in diagnosing,
CC
     treating or preventing NOVX-associated disorders, e.g. cancer, cachexia,
CC
     Alzheimer's disease, Parkinson's disease, obesity, diabetes, infectious
CC
     diseases, metabolic syndrome X or dyslipidaemias. The nucleic acids are
     further used as hybridisation probes, in chromosome mapping, tissue
CC
CC
     typing, preventive medicine, and pharmacogenomics. The present sequence
CC
     encodes a NOVX polypeptide of the invention.
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Query Match

Sequence 2881 BP; 527 A; 985 C; 867 G; 502 T; 0 U; 0 Other;

Score 2674.8; DB 12; Length 2881;

97.2%;

Best Local Similarity 98.9%; Pred. No. 0; 0; Matches 2727; Conservative Mismatches 22; Indels 1 CCGCGGGCCCCGCCCGCCCGCCCGCCGCCGCGCCATGGCCGTCCGGCCC 60 Qy Db 42 CCGCGGGCCCCGCCCGCCCGCCCGCCCGCCGCGCCATGGCCGTCCGGCCC 101 61 GGCCTGTGGCCAGCGCTCCTGGGCATAGTCCTCGCCGCTTGGCTCCGCGGCTCGGGTGCC 120 Qy . 102 GGCCTGTGGCCAGCGCTCCTGGGCATAGTCCTCGCCGCTTGGCTCCGCGGCTCGGGTGCC 161 Db 121 CAGCAGAGTGCCACCGTGGCCAACCCAGTGCCTGGTGCCAACCCGGACCTGCTTCCCCAC 180 Qу 162 CAGCAGAGTGCCACCGTGGCCAACCCAGTGCCTGGTGCCAACCCGGACCTGCTTCCCCAC 221 Db 181 TTCCTGGTGGAGCCCGAGGATGTGTACATCGTCAAGAACAAGCCAGTGCTGCTTGTGTGC 240 Qу 222 TTCCTGGTGGAGCCCGAGGATGTGTACATCGTCAAGAACAAGCCAGTGCTGCTTGTGTGC 281 Db 241 AAGGCCGTGCCCGCCACGCAGATCTTCTTCAAGTGCAACGGGGAGTGGGTGCGCCAGGTG 300 Qу 282 AAGGCCGTGCCCGCCACGCAGATCTTCTTCAAGTGCAACGGGGAGTGGGTGCGCCAGGTG 341 Db 301 GACCACGTGATCGAGCGCAGCACAGACGGGAGCAGTGGGCTGCCCACCATGGAGGTCCGC 360 Qy 342 GACCACGTGATCGAGCGCAGCACAGACGGGAGCAGTGGTGAGCCGACCATGGAGGTCCGC 401 Db 361 ATTAATGTCTCAAGGCAGCAGGTCGAGAAGGTGTTCGGGCTGGAGGAATACTGGTGCCAG 420 Qу 402 ATTAATGTCTCAAGGCAGCAGGTCGAGAAGGTGTTCGGGCTGGAGGAATACTGGTGCCAG 461 Db 421 TGCGTGGCATGGAGCTCCTCGGGCACCACAAGAGTCAGAAGGCCTACATCCGCATAGCC 480 Qу 462 TGCGTGGCATGGAGCTCCTCGGGCACCACAAGAGTCAGAAGGCCTACATCCGCATAGCC 521 Db 481 AGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTGTCCCTGGAGCAGGGC 540 Qу 522 AGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTGTCCCTGGAGCAGGGC 581 Db 541 ATCGTGCTGCCTTCCACCGGAGGGCATCCCTCCAGCCGAGGTGGAGTGGCTCCGG 600 Qу 582 ATCGTGCTGCCTCCACCGGAGGGCATCCCTCCAGCCGAGGTGGAGTGGCTCCGG 641 Db 601 AACGAGGACCTGGTGGACCCGTCCCTGGACCCCAATGTATACATCACGCGGGAGCACAGC 660 Qу 642 AACGAGGACCTGGTGGACCCGTCCCTGGACCCCAATGTATACATCACGCGGGAGCACAGC 701 Db 661 CTGGTGGTGCGACAGGCCCGCCTTGCTGACACGGCCAACTACACCTGCGTGGCCAAGAAC 720 Qу 702 CTGGTGGTGCGACAGGCCCGCCTTGCTGACACGGCCAACTACACCTGCGTGGCCAAGAAC 761 Db Qу 

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    New NOVX polypeptides and nucleic acid molecules useful for preventing or
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    The invention relates to a novel isolated polypeptide (NOVX). A
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    antilipaemic activity, and may have a use in gene therapy, and as a
CC
    vaccine. The polypeptides are encoded by NOVX polynucleotides comprising
    any of the 303 fully defined nucleotide sequences given in the
CC
CC
    specification. The polypeptide is useful in the manufacture of a
CC
    medicament for treating a syndrome associated with a human disease. The
CC
    polypeptide, polynucleotide and antibody are useful in diagnosing,
CC
    treating or preventing NOVX-associated disorders, e.g. cancer, cachexia,
CC
    Alzheimer's disease, Parkinson's disease, obesity, diabetes, infectious
CC
    diseases, metabolic syndrome X or dyslipidaemias. The nucleic acids are
CC
    further used as hybridisation probes, in chromosome mapping, tissue
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    encodes a NOVX polypeptide of the invention.
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    New NOVX polypeptides and nucleic acid molecules useful for preventing or
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PT
    obesity, and in chromosome mapping, tissue typing or pharmacogenomics.
XX
PS
    Example 21; SEQ ID NO 537; 1880pp; English.
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    The invention relates to a novel isolated polypeptide (NOVX). A
    polypeptide of the invention has cytostatic, immunomodulator,
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    neuroprotective, nootropic, anorectic, antidiabetic, antimicrobial, and
CC
CC
    antilipaemic activity, and may have a use in gene therapy, and as a
CC
    vaccine. The polypeptides are encoded by NOVX polynucleotides comprising
CC
    any of the 303 fully defined nucleotide sequences given in the
    specification. The polypeptide is useful in the manufacture of a
CC
    medicament for treating a syndrome associated with a human disease. The
CC
    polypeptide, polynucleotide and antibody are useful in diagnosing,
CC
    treating or preventing NOVX-associated disorders, e.g. cancer, cachexia,
CC
CC
    Alzheimer's disease, Parkinson's disease, obesity, diabetes, infectious
CC
    diseases, metabolic syndrome X or dyslipidaemias. The nucleic acids are
CC
    further used as hybridisation probes, in chromosome mapping, tissue
CC
    typing, preventive medicine, and pharmacogenomics. The present sequence
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    encodes a NOVX polypeptide of the invention.
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	Qy	481	AGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTGTCCCTGGAGCAGGGC	540
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Qy :	2575	AGCCATCTCAGCTTCTTTGCCTCCAAGCCCAGCCCCACAGCCATGATCCTCAACCTGTGG 2634
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Qу	2695	GGCCAGCCAGACGCTGGCCTCTTCACAGTGTCGGAGGCTGAGTGCTGAGGCCGGCC
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ID ADH71629 standard; DNA; 2881 BP.

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AC ADH71629;

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XX
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     New NOVX polypeptides and nucleic acid molecules useful for preventing or
PT
     treating NOVX-associated disorders, e.g. cancer, diabetes, infection or
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     obesity, and in chromosome mapping, tissue typing or pharmacogenomics.
```

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XX
PS
    Example 21; SEQ ID NO 525; 1880pp; English.
XX
CC
    The invention relates to a novel isolated polypeptide (NOVX). A
    polypeptide of the invention has cytostatic, immunomodulator,
CC
CC
    neuroprotective, nootropic, anorectic, antidiabetic, antimicrobial, and
CC
    antilipaemic activity, and may have a use in gene therapy, and as a
CC
    vaccine. The polypeptides are encoded by NOVX polynucleotides comprising
    any of the 303 fully defined nucleotide sequences given in the
CC
CC
    specification. The polypeptide is useful in the manufacture of a
    medicament for treating a syndrome associated with a human disease. The
CC
    polypeptide, polynucleotide and antibody are useful in diagnosing,
CC
    treating or preventing NOVX-associated disorders, e.g. cancer, cachexia,
CC
CC
    Alzheimer's disease, Parkinson's disease, obesity, diabetes, infectious
    diseases, metabolic syndrome X or dyslipidaemias. The nucleic acids are
CC
    further used as hybridisation probes, in chromosome mapping, tissue
CC
    typing, preventive medicine, and pharmacogenomics. The present sequence
CC
    encodes a NOVX polypeptide of the invention.
CC
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    Sequence 2881 BP; 526 A; 986 C; 868 G; 501 T; 0 U; 0 Other;
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 Query Match
                      97.28;
                             Score 2674.8; DB 12; Length 2881;
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Qу

Db

Qу

Db

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Db	582	ATCGTGCTGCCGTCCACCGGAGGGCATCCCTCCAGCCGAGGTGGAGTGGCTCCGG	641
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Db	702	CTGGTGGTGCGACAGGCCCGCCTTGCTGACACGGCCAACTACACCTGCGTGGCCAAGAAC	761
Qy	721	ATCGTGGCACGTCGCCGCAGCGCCTCCGCTGTCATCGTCTACGTGAACGGTGGGTG	780
Db	762	ATCGTGGCACGTCGCCGCGCGCTCCGCTGCTGTCATCGTCTACGTGAACGGTGGGTG	821
Qy	781	TCGACGTGGACCGAGTGGTCCGTCTGCAGCGCCAGCTGTGGGCGCGGCTGGCAGAAACGG	840
Db	822	TCGACGTGGACCGAGTGGTCCGTCTGCAGCGCCAGCTGTGGGCCGGCTGGCAGAAACGG	881
Qy	841	AGCCGGAGCTGCACCAACCCGGCGCCTCTCAACGGGGGCGCTTTCTGTGAGGGGCAGAAT	900
Db	882	AGCCGGAGCTGCACCAACCCGGCGCCTCTCAACGGGGGCGCTTTCTGTGAGGGGCAGAAT	941
Qy	901	GTCCAGAAAACAGCCTGCGCCACCCTGTGCCCAGTAGACGGCAGCTGGAGCCCGTGG	957
Db	942	GTCCATGACCGCACCGTCTCTCTCTGCTTGTCTCTGTGGACGGCAGCTGGAGCCCGTGG	1001
Qу	958	AGCAAGTGGTCGGCCTGTGGGCTGGACTGCACCCACTGGCGGAGCCGTGAGTGCTCTGAC	1017
Db	1002	AGCAAGTGGTCGGCCTGTGGGCTGGACTGCACCCACTGGCGGAGCCGTGAGTGCTCTGAC	1061
Qу	1018	CCAGCACCCCGCAACGGAGGGAGGAGTGCCAGGGCACTGACCTGGACACCCGCAACTGT	1077
Db	1062	CCAGCACCCGCAACGGAGGGGAGGGGCACTGACCTGGACACCCGCAACTGT	1121
Qу	1078	ACCAGTGACCTCTGTGTACACAGTGCTTCTGGCCCTGAGGACGTGGCCCTCTATGTGGGC	1137
Db	1122	ACCAGTGACCTCTGTGTACACAGTGCTTCTGGCCCTGAGGACGTGGCCCTCTATGTGGGC	1181
Qу	1138	CTCATCGCCGTGGCCGTCTGCTGGTCCTGCTGCTGTTGTCCTCATCCTCGTTTATTGC	1197
Db	1182	CTCATCGCCGTGGCCGTCTGCCTGGTCCTGCTGCTGCTTGTCCCCATCCTCGTTTATTGC	1241
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Db	1302	CAGCCCGTCAGCATCAAGCCCAGCAAAGCAGACAACCCCCATCTGCTCACCATCCAGCCG	1361
Ov	1318	GACTTCAGCACCACCACCACCTACCAGGGCAGTCTCTGTCCCCGGCAGGATGGGCCC	1377

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Db	1362	GACCTCAGCACCACCACCACCTACCAGGGCAGTCTCTGTCCCCGGCAGGATGGGCCC	1418
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Qу	1438	ACACTGCACCACGCTCTCAGGCCGAGGAGTTCGTCTCCCGCCTCTCCACC	1497
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Qу	1798	GACAGCTGGAGCCTGCGCCTCAAAAAGCAGTCGTGCGAGGGCAGCTGGGAGGATGTG	1854
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Qy	1855	CTGCACCTGGGCGAGGAGGCCCCTCCCACCTCTACTACTGCCAGCTGGAGGCCAGTGCC	1914
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Qy	1915	TGCTACGTCTTCACCGAGCAGCTGGGCCGCTTTGCCCTGGTGGGAGAGGCCCTCAGCGTG	1974
Db	1959	TGCTACGTCTTCACCGAGCAGCTGGGCCGCTTTGCCCTGGTGGGAGAGGCCCTCAGCGTG	2018
Qy	1975	GCTGCCGCCAAGCGCCTCAAGCTGCTTCTGTTTGCGCCGGTGGCCTGCACCTCCCTC	2034
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Qу	2035	TACAACATCCGGGTCTACTGCCTGCATGACACCCACGATGCACTCAAGGAGGTGGTGCAG	2094
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   obesity; diabetes; infectious disease; metabolic syndrome X;
KW
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   dyslipidaemia.
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Homo sapiens.

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DR
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     P-PSDB; ADH71632.
XX
    New NOVX polypeptides and nucleic acid molecules useful for preventing or
PT
     treating NOVX-associated disorders, e.g. cancer, diabetes, infection or
PT
     obesity, and in chromosome mapping, tissue typing or pharmacogenomics.
PT
XX
PS
     Example 21; SEQ ID NO 527; 1880pp; English.
XX
     The invention relates to a novel isolated polypeptide (NOVX). A
CC
CC
     polypeptide of the invention has cytostatic, immunomodulator,
CC
     neuroprotective, nootropic, anorectic, antidiabetic, antimicrobial, and
     antilipaemic activity, and may have a use in gene therapy, and as a
CC
     vaccine. The polypeptides are encoded by NOVX polynucleotides comprising
CC
     any of the 303 fully defined nucleotide sequences given in the
CC
CC
     specification. The polypeptide is useful in the manufacture of a
     medicament for treating a syndrome associated with a human disease. The
CC
CC
     polypeptide, polynucleotide and antibody are useful in diagnosing,
CC
     treating or preventing NOVX-associated disorders, e.g. cancer, cachexia,
```

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Alzheimer's disease, Parkinson's disease, obesity, diabetes, infectious
   diseases, metabolic syndrome X or dyslipidaemias. The nucleic acids are
CC
CC
    further used as hybridisation probes, in chromosome mapping, tissue
   typing, preventive medicine, and pharmacogenomics. The present sequence
CC
   encodes a NOVX polypeptide of the invention.
CC
XX
   Sequence 2881 BP; 526 A; 984 C; 868 G; 503 T; 0 U; 0 Other;
SO
                    97.2%;
                          Score 2674.8;
                                     DB 12;
                                            Length 2881;
 Query Match
                    98.9%;
 Best Local Similarity
                          Pred. No. 0:
 Matches 2727; Conservative
                            Mismatches
                                          Indels
                                                  9;
                                                            3;
                         0;
                                      22;
                                                     Gaps
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Qy
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Qу
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Qу
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Qу
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Db
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Qу
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Qy
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Db
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Qу

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Qу	721	ATCGTGGCACGTCGCCGCAGCGCCTCCGCTGTCATCGTCTACGTGAACGGTGGGTG	780
Db	762		821
Qу	781	TCGACGTGGACCGAGTGGTCCGTCTGCAGCGCCAGCTGTGGGCGCGGCTGGCAGAAACGG	840
Db	822	TCGACGTGGACCGAGTGGTCCGTCTGCAGCGCCAGCTGTGGGCGCGGCTGGCAGAAACGG	881
Qу	841	AGCCGGAGCTGCACCAACCCGGCGCCTCTCAACGGGGGCGCTTTCTGTGAGGGGCAGAAT	900
Db	882	AGCCGGAGCTGCACCAACCCGGCGCCTCTCAACGGGGGCGCTTTCTGTGAGGGGCAGAAT	941
Qу	901	GTCCAGAAAACAGCCTGCGCCACCCTGTGCCCAGTAGACGGCAGCTGGAGCCCGTGG	957
Db	942	GTCCATGACCGCACCGTCTCCTCTCTGCTTGTCTCTGTGGACGGCAGCTGGAGCCCGTGG	1001
Qу	958	AGCAAGTGGTCGGCCTGTGGGCTGGACTGCACCCACTGGCGGAGCCGTGAGTGCTCTGAC	1017
Db	1002	AGCAAGTGGTCGGCCTGTGGGCTGGACTGCACCCACTGGCGGAGCCGTGAGTGCTCTGAC	1061
Qу	1018	CCAGCACCCGCAACGGAGGGAGGGGAGTGCCAGGGCACTGACCTGGACACCCGCAACTGT	1077
Db	1062	CCAGCACCCGCAACGGAGGGAGGAGTGCCAGGGCACTGACCTGGACACCCGCAACTGT	1121
Qу	1078	ACCAGTGACCTCTGTGTACACAGTGCTTCTGGCCCTGAGGACGTGGCCCTCTATGTGGGC	1137
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Qу	1138	CTCATCGCCGTGGCCGTCTGCCTGGTCCTGCTGCTGCTTGTCCTCATCCTCGTTTATTGC	1197
Db	1182	CTCATCGCCGTGGCCGTCTGCCTGCTGCTGCTGCTCTCATCCTCGTTTATTGC	1241
Qу	1198	CGGAAGAAGGAGGGCTGGACTCAGATGTGGCTGACTCGTCCATTCTCACCTCAGGCTTC	1257
Db	1242	CGGAAGAAGGAGGGCTGGACTCAGATGTGGCTGACTCGTCCATTCTCACCTCAGGCTTC	1301
Qу	1258	CAGCCCGTCAGCATCAAGCCCAGCAAAGCAGACAACCCCCATCTGCTCACCATCCAGCCG	1317
Db	1302	CAGCCGTCAGCATCAAGCCCAGCAAAGCAGACAACCCCCATCTGCTCACCATCCAGCCG	1361
Qу	1318	GACCTCAGCACCACCACCACCTACCAGGGCAGTCTCTGTCCCCGGCAGGATGGGCCC	1377
Db	1362	GACCTCAGCACCACCACCACCTACCAGGGCAGTCTCTGTCCCCGGCAGGATGGGCCC	1418
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Qу	1438	ACACTGCACCACAGCTCTCCACCTCTGAGGCCGAGGAGTTCGTCTCCCGCCTCTCCACC	149.7
ĎР	1479	ACACTGCACCACACCTCTGAGGCCGAGGAGTTCGTCTCCCGCCTCTCCACC	1538
Qу	1498	CAGAACTACTTCCGCTCCCTGCCCCGAGGCACCAGCAACATGACCTATGGGACCTTCAAC	1557

Db	1539	CAGAACTACTTCCGCTCCCTGCCCCGAGGCACCAGCAACATGACCTATGGGACCTTCAAC	1598
Qy	1558	TTCCTCGGGGGCCGGCTGATGATCCCTAATACAGGTATCAGCCTCCTCATCCCCCAGAT	1617
Db	1599	TTCCTCGGGGGCCGGCTGATGATCCCTAATACAGGTATCAGCCTCCTCATCCCCCAGAT	1658
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Db	1659	GTCATACCCCGAGGGAAGATCTATGAGATCTACCTCACGCTGCACAAGCCGGAAGACGTG	1718
Qy	1678	AGGTTGCCCCTAGCTGGCTGTCAGACCCTGCTGAGTCCCATCGTTAGCTGTGGACCCCCT	1737
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Qу	1798	GACAGCTGGAGCCTGCGCCTCAAAAAGCAGTCGTGCGAGGGCAGCTGGGAGGATGTG	1854
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Qу	1855	CTGCACCTGGGCGAGGAGGCGCCCTCCCACCTCTACTACTGCCAGCTGGAGGCCAGTGCC	1914
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Qу	1975	GCTGCCGCCAAGCCCTCAAGCTGCTTCTGTTTGCGCCGGTGGCCTGCACCTCCCTC	2034
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     New NOVX polypeptides and nucleic acid molecules useful for preventing or
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     The invention relates to a novel isolated polypeptide (NOVX). A
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     antilipaemic activity, and may have a use in gene therapy, and as a
CC
     vaccine. The polypeptides are encoded by NOVX polynucleotides comprising
CC
     any of the 303 fully defined nucleotide sequences given in the
CC
     specification. The polypeptide is useful in the manufacture of a
CC
     medicament for treating a syndrome associated with a human disease. The
CC
     polypeptide, polynucleotide and antibody are useful in diagnosing,
CC
     treating or preventing NOVX-associated disorders, e.g. cancer, cachexia,
CC
     Alzheimer's disease, Parkinson's disease, obesity, diabetes, infectious
CC
     diseases, metabolic syndrome X or dyslipidaemias. The nucleic acids are
CC
     further used as hybridisation probes, in chromosome mapping, tissue
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     typing, preventive medicine, and pharmacogenomics. The present sequence
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-	QУ	601	AACGAGGACCTGGTGGACCCGTCCCTGGACCCCAATGTATACATCACGCGGGAGCACAGC	660
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    Alsobrook JP, Alvarez E, Anderson DW, Boldog FL, Casman SJ;
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    Catterton E, Chapoval A, Crabtree-Bokor JR, Edinger SR, Ellerman K;
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    Ettenberg S, Gangolli EA, Gerlach VL, Gorman L, Gunther E, Guo X;
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    Gusev VY, Herrmann JL, Ji W, Kekuda R, Li L, Liu X, Macdougall JR;
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    Maclachlan T, Malyankar UM, Mezick AJ, Millet I, Mishra VS;
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    Padigaru M, Patturajan M, Pena CEA, Peyman JA, Raha D, Rastelli L; Rieger DK, Rothenberg ME, Sciore P, Shenoy SG, Shimkets RA;
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    Smithson G, Spytek KA, Stone DJ, Vernet CAM, Voss EZ, Zhong M;
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    Zhong H;
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    WPI; 2004-081935/08.
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    P-PSDB; ADH71628.
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    New NOVX polypeptides and nucleic acid molecules useful for preventing or
    treating NOVX-associated disorders, e.g. cancer, diabetes, infection or
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PT
    obesity, and in chromosome mapping, tissue typing or pharmacogenomics.
XX
PS
    Example 21; SEQ ID NO 523; 1880pp; English.
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CC
    The invention relates to a novel isolated polypeptide (NOVX). A
CC
    polypeptide of the invention has cytostatic, immunomodulator,
    neuroprotective, nootropic, anorectic, antidiabetic, antimicrobial, and
CC
CC
    antilipaemic activity, and may have a use in gene therapy, and as a
    vaccine. The polypeptides are encoded by NOVX polynucleotides comprising
CC
CC
    any of the 303 fully defined nucleotide sequences given in the
    specification. The polypeptide is useful in the manufacture of a
CC
CC
    medicament for treating a syndrome associated with a human disease. The
    polypeptide, polynucleotide and antibody are useful in diagnosing,
CC
CC
    treating or preventing NOVX-associated disorders, e.g. cancer, cachexia,
    Alzheimer's disease, Parkinson's disease, obesity, diabetes, infectious
CC
CC
    diseases, metabolic syndrome X or dyslipidaemias. The nucleic acids are
CC
    further used as hybridisation probes, in chromosome mapping, tissue
CC
    typing, preventive medicine, and pharmacogenomics. The present sequence
CC
    encodes a NOVX polypeptide of the invention.
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Qy	361	ATTAATGTCTCAAGGCAGCAGGTCGAGAAGGTGTTCGGGCTGGAGGAATACTGGTGCCAG	420
Db	402	ATTAATGTCTCAAGGCAGCAGGTCGAGAAGGTGTTCGGGCTGGAGGAATACTGGTGCCAG	461
Qy	421	TGCGTGGCATGGAGCTCCTCGGGCACCACCAAGAGTCAGAAGGCCTACATCCGCATAGCC	480
Db	462	TGCGTGGCATGGAGCTCCTCGGGCACCACCAAGAGTCAGAAGGCCTACATCCGCATAGCC	521
Qу	481	AGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTGTCCCTGGAGCAGGGC	540
Db	522	AGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTGTCCCTGGAGCAGGGC	581
Qу	541	ATCGTGCTGCCCTCCACCGGAGGGCATCCCTCCAGCCGAGGTGGATGGCTCCGG	600
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Qy	601	AACGAGGACCTGGTGGACCCGTCCCTGGACCCCAATGTATACATCACGCGGGAGCACAGC	660
Db	642	AACGAGGACCTGGTGGACCCCTGGACCCCAATGTATACATCACGCGGGAGCACAGC	701
Qy	661	CTGGTGGTGCGACAGGCCCGCCTTGCTGACACGGCCAACTACACCTGCGTGGCCAAGAAC	720
Db	702	CTGGTGGTGCGACAGGCCCGCCTTGCTGACACGGCCAACTACACCTGCGTGGCCAAGAAC	761
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Qy .	781	TCGACGTGGACCGAGTGGTCCGTCTGCAGCGCCAGCTGTGGGCGGCGGCTGGCAGAAACGG	840
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Qу	901	GTCCAGAAAACAGCCTGCGCCACCCTGTGCCCAGTAGACGGCAGCTGGAGCCCGTGG	957
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Qу	1018	CCAGCACCCGCAACGGAGGGAGGGGAGTGCCAGGGCACTGACCTGGACACCCGCAACTGT	1077
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Qу	1258	CAGCCCGTCAGCATCAAGCCCAGCAAAGCAGACAACCCCCATCTGCTCACCATCCAGCCG	1317
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Qу	1378	AGCCCCAAGTTCCAGCTCACCAATGGGCACCTGCTCAGCCCCCTGGGTGGCGGCCGCCAC	1437
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     vaccine; cancer; cachexia; Alzheimer's disease; Parkinson's disease;
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ΡI
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ΡI
     Smithson G, Spytek KA, Stone DJ, Vernet CAM, Voss EZ, Zhong M;
ΡI
PΙ
     Zhong H;
XX
DR
     WPI; 2004-081935/08.
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DR
    P-PSDB; ADH71640.
XX
    New NOVX polypeptides and nucleic acid molecules useful for preventing or
PT
    treating NOVX-associated disorders, e.g. cancer, diabetes, infection or
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    obesity, and in chromosome mapping, tissue typing or pharmacogenomics.
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XX
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    Example 21; SEQ ID NO 535; 1880pp; English.
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CC
    polypeptide of the invention has cytostatic, immunomodulator,
    neuroprotective, nootropic, anorectic, antidiabetic, antimicrobial, and
CC
    antilipaemic activity, and may have a use in gene therapy, and as a
CC
CC
    vaccine. The polypeptides are encoded by NOVX polynucleotides comprising
CC
    any of the 303 fully defined nucleotide sequences given in the
    specification. The polypeptide is useful in the manufacture of a
CC
CC
    medicament for treating a syndrome associated with a human disease. The
    polypeptide, polynucleotide and antibody are useful in diagnosing,
CC
    treating or preventing NOVX-associated disorders, e.g. cancer, cachexia,
CC
    Alzheimer's disease, Parkinson's disease, obesity, diabetes, infectious
CC
    diseases, metabolic syndrome X or dyslipidaemias. The nucleic acids are
CC
CC
    further used as hybridisation probes, in chromosome mapping, tissue
    typing, preventive medicine, and pharmacogenomics. The present sequence
CC
    encodes a NOVX polypeptide of the invention.
CC
XX
    Sequence 2881 BP; 525 A; 985 C; 869 G; 502 T; 0 U; 0 Other;
SQ
 Query Match
                      97.2%;
                            Score 2674.8; DB 12; Length 2881;
 Best Local Similarity
                      98.9%; Pred. No. 0;
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 Matches 2727; Conservative
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Qу	481	AGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTGTCCCTGGAGCAGGGC	540
Db	522	AGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTGTCCCTGGAGCAGGGC	581
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Qy	1078	ACCAGTGACCTCTGTGTACACAGTGCTTCTGGCCCTGAGGACGTGGCCCTCTATGTGGGC	1137
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Qу	1138	CTCATCGCCGTGGCCGTCTGCCTGGTCCTGCTGCTGCTCATCCTCGTTTATTGC	1197
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Db	1242		1301

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Qу	1258	CAGCCCGTCAGCATCAAGCCCAGCAAAGCAGCAACCCCCATCTGCTCACCATCCAGCCG	1317	
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vaccine; cancer; cachexia; Alzheimer's disease; Parkinson's disease;

KW

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                                                                 Rastelli L;
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     Rieger DK, Rothenberg ME,
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DR
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XX
PT
    New NOVX polypeptides and nucleic acid molecules useful for preventing or
PT
     treating NOVX-associated disorders, e.g. cancer, diabetes, infection or
PT
     obesity, and in chromosome mapping, tissue typing or pharmacogenomics.
XX
PS
     Example 21; SEQ ID NO 539; 1880pp; English.
XX
CC
     The invention relates to a novel isolated polypeptide (NOVX). A
CC
     polypeptide of the invention has cytostatic, immunomodulator,
CC
     neuroprotective, nootropic, anorectic, antidiabetic, antimicrobial, and
CC
     antilipaemic activity, and may have a use in gene therapy, and as a
CC
     vaccine. The polypeptides are encoded by NOVX polynucleotides comprising
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CC any of the 303 fully defined nucleotide sequences given in the specification. The polypeptide is useful in the manufacture of a CC CC medicament for treating a syndrome associated with a human disease. The CC polypeptide, polynucleotide and antibody are useful in diagnosing, treating or preventing NOVX-associated disorders, e.g. cancer, cachexia, CC CC Alzheimer's disease, Parkinson's disease, obesity, diabetes, infectious diseases, metabolic syndrome X or dyslipidaemias. The nucleic acids are CC CC further used as hybridisation probes, in chromosome mapping, tissue CC typing, preventive medicine, and pharmacogenomics. The present sequence CC encodes a NOVX polypeptide of the invention. XX

SQ

Sequence 2881 BP; 527 A; 985 C; 867 G; 502 T; 0 U; 0 Other;

Query Match 97.2%; Score 2674.8; DB 12; Length 2881; Best Local Similarity 98.9%; Pred. No. 0; Matches 2727; Conservative 0; Mismatches 22; Indels 9; Gaps 3;

	Matches	212	, conservative o, rismacches 22, finders o, daps	,
Q	!.	1	CCGCGGGCCCGCCCGCCCGCCCGCCCGCCCGCCCATGGCCGTCCGGCCC 60	
Dk	)	42	CCGCGGGGCCCGCCCGCCCGCCCGCCCGCCCGCCCGCC	1
Qy	7	61	GGCCTGTGGCCAGCGCTCCTGGGCATAGTCCTCGCCGCTTGGCTCCGCGGCTCGGGTGCC 120	C
Dk	)	102	GGCCTGTGGCCAGCGCTCCTGGGCATAGTCCTCGCCGCTTGGCTCCGCGGCTCGGGTGCC 161	1
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Dk	<b>)</b>	162	CAGCAGAGTGCCACCGTGGCCAACCCAGTGCCTGGTGCCAACCCGGACCTGCTTCCCCAC 221	1
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Dk	>	222	TTCCTGGTGGAGCCCGAGGATGTGTACATCGTCAAGAACAAGCCAGTGCTGCTTGTGTGC 281	1
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Dk		282	AAGGCCGTGCCCGCCACGCAGATCTTCTTCAAGTGCAACGGGGAGTGGGTGCGCCAGGTG 341	1
Qy	7	301	GACCACGTGATCGAGCGCAGCACAGACGGGAGCAGTGGGCTGCCCACCATGGAGGTCCGC 360	Э
Dk		342	GACCACGTGATCGAGCGCAGCACAGACGGGAGCAGTGGTGAGCCGACCATGGAGGTCCGC 401	1
Qy	7	361	ATTAATGTCTCAAGGCAGCAGGTCGAGAAGGTGTTCGGGCTGGAGGAATACTGGTGCCAG 420	Э
Dł	)	402	ATTAATGTCTCAAGGCAGCAGGTCGAGAAGGTGTTCGGGCTGGAGGAATACTGGTGCCAG 461	1
Qy	7	421	TGCGTGGCATGGAGCTCCTCGGGCACCACCAAGAGTCAGAAGGCCTACATCCGCATAGCC 480	Э
Dł		462	TGCGTGGCATGGAGCTCCTCGGGCACCACCAAGAGTCAGAAGGCCTACATCCGCATAGCC 521	1
Q	7	481	AGATTGCGCAAGAACTTCGAGCAGGAGCCGCTGGCCAAGGAGGTGTCCCTGGAGCAGGGC 540	0
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Q	7	541	ATCGTGCTGCCCTCCACCGGAGGGCATCCCTCCAGCCGAGGTGGAGTGGCTCCGG 600	0
D}	<b>)</b>	582	ATCGTGCTGCCCTCCACCGGAGGGCATCCCTCCAGCCGAGGTGGAGTGGCTCCGG 64	1

Qу	601	AACGAGGACCTGGTGGACCCGTCCCTGGACCCCAATGTATACATCACGCGGGAGCACAGC	660
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Qу	721	ATCGTGGCACGTCGCCGCAGCGCCTCCGCTGTCATCGTCTACGTGAACGGTGGGTG	780
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Db ·	1302	CAGCCCGTCAGCATCAAGCCCAGCAAAGCAGACAACCCCCATCTGCTCACCATCCAGCCG	1361
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Qy	1378	AGCCCCAAGTTCCAGCTCACCAATGGGCACCTGCTCAGCCCCCTGGGTGGCGGCCGCCAC	1437
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Qy	1558	TTCCTCGGGGGCCGGCTGATGATCCCTAATACAGGTATCAGCCTCCTCATCCCCCAGAT	1617
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Db	1779	GGCGTCCTGCTCACCCGGCCAGTCATCCTGGCTATGGACCACTGTGGGGAGCCCAGCCCT	1838
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Qу	1855	CTGCACCTGGGCGAGGAGGCGCCCTCCCACCTCTACTACTGCCAGCTGGAGGCCAGTGCC	1914
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## GenCore version 5.1.6 Copyright (c) 1993 - 2005 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: March 5, 2005, 19:26:23; Search time 433.33 Seconds

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10391.692 Million cell updates/sec

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Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0

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Post-processing: Minimum Match 0%

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Listing first 45 summaries

Database: Issued Patents NA:\*

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

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Ţ	2259	82.1	3014	2	US-08-808-982-1	Sequence 1, Appli
2	2259	82.1	3014	3	US-09-306-902A-1	Sequence 1, Appli
3	1562.4	56.8	1787	2	US-08-808-982-2	Sequence 2, Appli
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7	833.6	30.3	3008	4	US-09-949-016-4794	Sequence 4794, Ap
8	487	17.7	2736	4	US-09-969-532-9	Sequence 9, Appli
9	487	17.7	3411	4	US-09-969-532-33	Sequence 33, Appl
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## ALIGNMENTS

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RESULT 1
US-08-808-982-1
; Sequence 1, Application US/08808982
; Patent No. 5939271
  GENERAL INFORMATION:
    APPLICANT: Tessier-Lavigne, Marc
    APPLICANT: Leonardo, E. David
    APPLICANT: Hink, Lindsay
    APPLICANT: Masu, Masayuki
    APPLICANT: Kazuko, Keino-Masu
    TITLE OF INVENTION: Netrin Receptors
    NUMBER OF SEQUENCES: 8
    CORRESPONDENCE ADDRESS:
      ADDRESSEE: SCIENCE & TECHNOLOGY LAW GROUP
      STREET: 268 BUSH STREET, SUITE 3200
      CITY: SAN FRANCISCO
      STATE: CALIFORNIA
;
      COUNTRY: USA
```

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ZIP: 94104
   COMPUTER READABLE FORM:
     MEDIUM TYPE: Floppy disk
     COMPUTER: IBM PC compatible
     OPERATING SYSTEM: PC-DOS/MS-DOS
     SOFTWARE: PatentIn Release #1.0, Version #1.30
   CURRENT APPLICATION DATA:
     APPLICATION NUMBER: US/08/808,982
     FILING DATE:
     CLASSIFICATION: 530
   ATTORNEY/AGENT INFORMATION:
     NAME: OSMAN, RICHARD A
     REGISTRATION NUMBER: 36,627
     REFERENCE/DOCKET NUMBER: UC96-217
   TELECOMMUNICATION INFORMATION:
     TELEPHONE: (415) 343-4341
     TELEFAX: (415) 343-4342
  INFORMATION FOR SEQ ID NO: 1:
   SEQUENCE CHARACTERISTICS:
     LENGTH: 3014 base pairs
     TYPE: nucleic acid
     STRANDEDNESS: double
     TOPOLOGY: linear
   MOLECULE TYPE: cDNA
US-08-808-982-1
 Query Match
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 Best Local Similarity 89.7%; Pred. No. 0;
 Matches 2427; Conservative
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